(19) World Intellectual Property Organization International Bureau





(43) International Publication Date 9 August 2001 (09.08.2001)

PCT

(10) International Publication Number WO 01/57277 A2

(51) International Patent Classification⁷: C07K 14/47, C07H 21/04

C12Q 1/68,

(21) International Application Number: PCT/US01/00669

(22) International Filing Date: 30 January 2001 (30.01.2001)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

60/180;312	4 February 2000 (04.02.2000)	US
60/207,456	26 May 2000 (26.05.2000)	US
09/608,408	30 June 2000 (30.06.2000)	US
09/632,366	3 August 2000 (03.08.2000)	US
60/234,687	21 September 2000 (21.09.2000)	US
60/236,359	27 September 2000 (27.09.2000)	US
0024263.6	4 October 2000 (04.10.2000)	GB

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- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

 without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

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(54) Title: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR ANALYSIS OF GENE EXPRESSION IN HUMAN FETAL LIVER

(57) Abstract: A single exon nucleic acid microarray comprising a plurality of single exon nucleic acid probes for measuring gene expression in a sample derived from human Fetal liver is described. Also described are single exon nucleic acid probes expressed in the Fetal liver and their use in methods for detecting gene expression.

HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR ANALYSIS OF GENE EXPRESSION IN HUMAN FETAL LIVER

CROSS REFERENCE TO RELATED APPLICATIONS

The present application is a continuation-in-part of U.S. patent application serial nos. 09/632,366, filed August 3, 2000 and 09/608,408, filed June 30, 2000; claims the benefit under 35 U.S.C. s 119(e) of U.S.provisional patent application serial nos. 60/236,359, filed September 27, 2000, 60/234,687, filed September 21, 2000, 60/207,456, filed May 26, 2000, and 60/180,312, filed February 4, 2000; and further claims the benefit under 35 U.S.C. s 119(a) of UK patent application no. 0024263.6, filed October 4, 2000, the disclosures of which are incorporated herein by reference in their entireties.

REFERENCE TO SEQUENCE LISTING AND INCORPORATION BY REFERENCE THEREOF

The present application includes a Sequence Listing in electronic format, filed pursuant to PCT Administrative Instructions 801 - 806 on a single CD-R disc, in triplicate, containing a file named pto_FETAL_LIVER.txt, created 24 January 2001, having 25,630,231 bytes. The Sequence Listing contained in said file on said disc is incorporated herein by reference in its entirety.

Field of the Invention

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The present invention relates to genome-derived single exon microarrays useful for verifying the expression of regions of genomic DNA predicted to encode protein. In particular, the present invention relates to unique genomederived single exon nucleic acid probes expressed in human

Fetal liver and single exon nucleic acid microarrays that include such probes.

Background of the Invention

For almost two decades following the invention of general techniques for nucleic acid sequencing, Sanger et al., Proc. Natl. Acad. Sci. USA 70(4):1209-13 (1973); Gilbert et al., Proc. Natl. Acad. Sci. USA 70(12):3581-4 (1973), these techniques were used principally as tools to further the understanding of proteins — known or suspected — about which a basic foundation of biological knowledge had already been built. In many cases, the cloning effort that preceded sequence identification had been both informed and directed by that antecedent biological understanding.

For example, the cloning of the T cell receptor for antigen was predicated upon its known or suspected cell type-specific expression, by its suspected membrane association, and by the predicted assembly of its gene via T cell-specific somatic recombination. Subsequent sequencing efforts at once confirmed and extended understanding of this family of proteins. Hedrick et al., Nature 308(5955):153-8 (1984).

More recently, however, the development of high
throughput sequencing methods and devices, in concert with
large public and private undertakings to sequence the human
and other genomes, has altered this investigational
paradigm: today, sequence information often precedes
understanding of the basic biology of the encoded protein
product.

One of the approaches to large-scale sequencing is predicated upon the proposition that expressed sequences — that is, those accessible through isolation of mRNA — are of greatest initial interest. This "expressed sequence tag" ("EST") approach has already yielded vast

amounts of sequence data (see for example Adams et al.,

Science 252:1651 (1991); Williamson, Drug Discov. Today

4:115 (1999)). For nucleic acids sequenced by this
approach, often the only biological information that is
known a priori with any certainty is the likelihood of
biologic expression itself. By virtue of the species and
tissue from which the mRNA had originally been obtained,
most such sequences are also annotated with the identity of
the species and at least one tissue in which expression
appears likely.

More recently, the pace of genomic sequencing has accelerated dramatically. When genomic DNA serves as the initial substrate for sequencing efforts, expression cannot be presumed; often the only a priori biological information about the sequence includes the species and chromosome (and perhaps chromosomal map location) of origin.

With the ever-accelerating pace of sequence accumulation by directed, EST, and genomic sequencing approaches — and in particular, with the accumulation of sequence information from multiple genera, from multiple species within genera, and from multiple individuals within a species — there is an increasing need for methods that rapidly and effectively permit the functions of nucleic sequences to be elucidated. And as such functional information accumulates, there is a further need for methods of storing such functional information in meaningful and useful relationship to the sequence itself; that is, there is an increasing need for means and apparatus for annotating raw sequence data with known or predicted functional information.

Although the increase in the pace of genomic sequencing is due in large part to technological changes in sequencing strategies and instrumentation, Service, Science 280:995 (1998); Pennisi, Science 283: 1822-1823 (1999), there is an important functional motivation as well.

While it was understood that the EST approach would rarely be able to yield sequence information about the noncoding portions of the genome, it now also appears the EST approach is capable of capturing only a fraction of a genome's actual expression complexity.

For example, when the C. elegans genome was fully sequenced, gene prediction algorithms identified over 19,000 potential genes, of which only 7,000 had been found by EST sequencing. C. elegans Sequencing Consortium, 10 Science 282:2012 (1998). Analogously, the recently completed sequence of chromosome 2 of Arabidopsis predicts over 4000 genes, Lin et al., Nature, 402:761 (1999), of which only about 6% had previously been identified via EST sequencing efforts. Although the human genome has the 15 greatest depth of EST coverage, it is still woefully short of surrendering all of its genes. One recent estimate suggests that the human genome contains more than 146,000 genes, which would at this point leave greater than half of the genes undiscovered. It is now predicted that many 20 genes, perhaps 20 to 50%, will only be found by genomic sequencing.

There is, therefore, a need for methods that permit the functional regions of genomic sequence — and most importantly, but not exclusively, regions that

25 function to encode genes — to be identified.

Much of the coding sequence of the human genome is not homologous to known genes, making detection of open reading frames ("ORFs") and predictions of gene function difficult. Computational methods exist for predicting coding regions in eukaryotic genomes. Gene prediction programs such as GRAIL and GRAIL II, Uberbacher et al., Proc. Natl. Acad. Sci. USA 88(24):11261-5 (1991); Xu et al., Genet. Eng. 16:241-53 (1994); Uberbacher et al., Methods Enzymol. 266:259-81 (1996); GENEFINDER, Solovyev et al., Nucl. Acids. Res. 22:5156-63 (1994); Solovyev et al.,

Ismb 5:294-302 (1997); and GENESCAN, Burge et al., J. Mol. Biol. 268:78-94 (1997), predict many putative genes without known homology or function. Such programs are known, however, to give high false positive rates. Burset et al., 5 Genomics 34:353-367 (1996). Using a consensus obtained by a plurality of such programs is known to increase the reliability of calling exons from genomic sequence. Ansari-Lari et al., Genome Res. 8(1):29-40 (1998)

Identification of functional genes from genomic 10 data remains, however, an imperfect art. For example, in reporting the full sequence of human chromosome 21, the Chromosome 21 Mapping and Sequencing Consortium reports that prior bioinformatic estimates of human gene number may need to be revised substantially downwards. Nature 15 405:311-199 (2000); Reeves, Nature 405:283-284 (2000).

Thus, there is a need for methods and apparatus that permit the functions of the regions identified bioinformatically - and specifically, that permit the expression of regions predicted to encode protein - readily 20 to be confirmed experimentally.

Recently, the development of nucleic acid microarrays has made possible the automated and highly parallel measurement of gene expression. Reviewed in Schena (ed.), DNA Microarrays : A Practical Approach (Practical Approach Series), Oxford University Press (1999) (ISBN: 0199637768); Nature Genet. 21(1)(suppl):1 - 60 (1999); Schena (ed.), Microarray Biochip: Tools and Technology, Eaton Publishing Company/BioTechniques Books Division (2000) (ISBN: 1881299376).

It is common for microarrays to be derived from cDNA/EST libraries, either from those previously described in the literature, such as those from the I.M.A.G.E. consortium, Lennon et al., Genomics 33(1):151-2 (1996), or from the construction of "problem specific" libraries 35 targeted at a particular biological question, R.S. Thomas

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PCT/US01/00669 WO 01/57277

et al., Cancer Res. (in press). Such microarrays by definition can measure expression only of those genes found in EST libraries, and thus have not been useful as probes for genes discovered solely by genomic sequencing.

. The utility of using whole genome nucleic acid microarrays to answer certain biological questions has been demonstrated for the yeast Saccharomyces cerevisiae. De Risi et al., Science 278:680 (1997). The vast majority of yeast nuclear genes, approximately 95% however, are single 10 exon genes, i.e., lack introns, Lopez et al., RNA 5:1135-1137 (1999); Goffeau et al., Science 274:563-67 (1996), permitting coding regions more readily to be identified. Whole genome nucleic acid microarrays have not generally been used to probe gene expression from more complex 15 eukaryotic genomes, and in particular from those averaging more than one intron per gene.

Diseases of the liver are a significant cause of human morbidity and mortality. Increasingly, genetic factors are being found that contribute to predisposition, 20 onset, and/or aggressiveness of most, if not all, of these diseases; although causative mutations in single genes have been identified for some, these disorders are believed for the most part to have polygenic etiologies. There is a need for methods and apparatus that permit prediction of 25 diseases of the liver, particularly those diseases with polygenic etiology, from diagnosis of fetal liver.

Summary of the Invention

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The present invention solves these and other problems in the art by providing methods and apparatus for predicting, confirming, and displaying functional information derived from genomic sequence. The present invention also provides apparatus for verifying the 35 expression of putative genes identified within genomic

sequence.

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In particular, the invention provides novel genome-derived single exon nucleic acid microarrays useful for verifying the expression of putative genes identified within genomic sequence.

The present invention also provides compositions and kits for the ready production of nucleic acids identical in sequence to, or substantially identical in sequence to, probes on the genome-derived single exon microarrays of the present invention.

Accordingly, in a first aspect of the invention, there is provided a spatially-addressable set of single exon nucleic acid probes for measuring gene expression in a sample derived from human Fetal liver, comprising a plurality of single exon nucleic acid probes according to any one of the nucleotide sequences set out in SEQ ID NOS: 1 - 12,673 or a complementary sequence, or a portion of such a sequence.

By plurality is meant at least two, suitably at least 20, most suitably at least 100, preferably at least 1000 and, most preferably, upto 5000.

In one embodiment of the first aspect, each of said plurality of probes is separately and addressably amplifiable.

In an alternative embodiment, each of said plurality of probes is separately and addressably isolatable from said plurality.

In a preferred embodiment, each of said plurality of probes is amplifiable using at least one common primer.

Preferably, each of said plurality of probes is amplifiable using a first and a second common primer.

In yet another embodiment, said set of single exon nucleic acid probes comprises between 50 - 20,000 probes, for example, 50 - 5000.

35 Suitably, said set of single exon nucleic acid

PCT/US01/00669 WO 01/57277

probes comprises at least 50 - 1000 discrete single exon nucleic acid probes having a sequence as set out in any of SEQ ID NOS.: 1 - 25,129 or a complimentary sequence, or a portion of such a sequence.

Preferably, the average length of the single exon nucleic acid probes is between 200 and 500 bp. It is preferred that the average length should be at least 200bp, suitably at least 250bp, most suitably at least 300bp, preferably at least 400bp and, most preferably, 500 bp.

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In another embodiment, the single exon nucleic acid probes lack prokaryotic and bacteriophage vector sequence. It is preferred that at least 50%, suitably at least 60%, most suitably at least 70%, preferably at least 75%, more preferably at least 80, 85, 90, 95 or 99% of said 15 single exon nucleic acid probes lack prokaryotic and bacteriophage vector sequence.

In another preferred embodiment, said single exon nucleic acid lack homopolymeric stretches of A or T. It is preferred that at least 50%, suitably at least 60%, most 20 suitably at least 70%, preferably at least 75%, more preferably at least 80, 85, 90, 95 or 99% of said single exon nucleic acid probes lack homopolymeric stretches of A or T.

Preferably, a spatially-addressable set of single 25 exon nucleic acid probes in accordance with the first aspect of the invention is is addressably disposed upon a substrate.

Suitable substrates include a filter membrane which may, preferably, be nitrocellulose or nylon. The 30 nylon may preferably, be positively-charged. Other suitable substrates include glass, amorphous silicon, crystalline silicon, and plastic. Further suitable materials include polymethylacrylic, polyethylene, polypropylene, polyacrylate, polymethylmethacrylate, polyvinylchloride, 35 polytetrafluoroethylene, polystyrene, polycarbonate,

polyacetal, polysulfone, celluloseacetate, cellulosenitrate, nitrocellulose, and mixtures thereof.

In a second aspect of the invention, there is provided a microarray comprising a spatially addressable set of single exon nucleic acid probes in accordance with the first aspect of the invention.

In one embodiment, a genome-derived single-exon microarray is packaged together with such an ordered set of amplifiable probes corresponding to the probes, or one or more subsets of probes, thereon. In alternative embodiments, the ordered set of amplifiable probes is packaged separately from the genome-derived single exon microarray.

In another aspect, the invention provides genomederived single exon nucleic acid probes useful for gene
expression analysis, and particularly for gene expression
analysis by microarray. In particular embodiments of this
aspect, the present invention provides human single-exon
probes that include specifically-hybridizable fragments of
SEQ ID Nos. 12,674 - 25,129, wherein the fragment
hybridizes at high stringency to an expressed human gene.
In particular embodiments, the invention provides single
exon probes comprising SEQ ID Nos. 1 - 12,673.

Accordingly, in a third aspect of the invention,

there is provided a single exon nucleic acid probe for
measuring human gene expression in a sample derived from
human Fetal liver which is a nucleic acid molecule
comprising a nucleotide sequence as set out in any of SEQ
ID NOs.: 1 - 12,673 or a complementary sequence or a

fragment thereof wherein said probe hybridizes at high
stringency to a nucleic acid expressed in the human Fetal
liver.

In one embodiment, a single exon nucleic acid probe in accordance with the third aspect comprises a nucleotide sequence as set out in any of SEQ ID NOs.:

PCT/US01/00669 WO 01/57277

12,674 - 25,129 or a complementary sequence or a fragment thereof.

In a fourth aspect of the invention, there is provided a single exon nucleic acid probe for measuring 5 human gene expression in a sample derived from human Fetal liver which is a nucleic acid molecule having a sequence encoding a peptide comprising a peptide sequence as set out in any of SEQ ID NOs.: 25,130 - 37,156 or a complementary sequence or a fragment thereof wherein said probe 10 hybridizes at high stringency to a nucleic acid expressed in the human Fetal liver.

Preferably, a single exon nucleic acid probe in accordance with the third or fourth aspects of the invention comprises between at least 15 and 50 contiguous 15 nucleotides of said SEQ ID NO:. It is preferred that the single exon nucleic acid probe comprises at least 15, suitably at least 20, more suitably at least 25 or preferably at least 50 contiguous nucleotides of said SEQ ID NO:.

In another preferred embodiment, a single exon nucleic acid probe in accordance with the third or fourth aspects of the invention is between 3kb and 25kb in length. It is preferred that said probe is no more than 3kb, suitably no more than 5kb, more suitably no more than 10kb, 25 preferably 15kb, more preferably 20kb or, most preferably, no more than 20kb in length.

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Preferably, a single exon nucleic acid probe in accordance with either the fifth or sixth aspect of the invention is DNA, preferably single-stranded DNA, RNA or PNA.

In another embodiment of either the third or fourth aspect of the invention, a single exon nucleic acid probe is detectably labeled. Suitable detectable labels include a radionuclide, a fluorescent label or a first 35 member of a specific binding pair. Suitable fluorescent

labels include dyes such as cyanine dyes, preferably Cy3 and Cy5 although other suitable dyes will be known to those skilled in the art.

In a particularly preferred embodiment, a single

sexon nucleic acid probe in accordance with either the third
or fourth aspect of the invention lacks prokaryotic and
bacteriophage vector sequence. In yet another embodiment, a
single exon nucleic acid probe in accordance with either
the third or fourth aspect of the invention lacks
homopolymeric stretches of A or T.

In a fifth aspect of the invention, there is provided an amplifiable nucleic acid composition, comprising:

the single exon nucleic acid probe in accordance
with either of the third or fourth aspects of the
invention; and at least one nucleic acid primer;

wherein said at least one primer is sufficient to prime enzymatic amplification of said probe.

In an sixth aspect of the invention, there is
provided a method of measuring gene expression in a sample
derived from human Fetal liver, comprising:

contacting the single exon microarray in accordance with the second aspect of the invention, with a first collection of detectably labeled nucleic acids, said first collection of nucleic acids derived from mRNA of human Fetal liver; and then

measuring the label detectably bound to each probe of said microarray.

In a seventh aspect of the invention, there is provided a method of identifying exons in a eukaryotic genome, comprising:

algorithmically predicting at least one exon from genomic sequence of said eukaryote; and then

detecting specific hybridization of detectably labeled nucleic acids to a single exon probe,

wherein said detectably labeled nucleic acids are derived from mRNA from the Fetal liver of said eukaryote, said probe is a single exon probe having a fragment identical in sequence to, or complementary in sequence to, said predicted exon, said probe is included within a single exon microarray in accordance with the first aspect of the invention, and said fragment is selectively hybridizable at high stringency.

In a eighth aspect of the invention, there is provided a method of assigning exons to a single gene, comprising:

identifying a plurality of exons from genomic sequence in accordance with the seventh aspect of the invention; and then

measuring the expression of each of said exons in a plurality of tissues and/or cell types using hybridization to single exon microarrays having a probe with said exon,

wherein a common pattern of expression of said 20 exons in said plurality of tissues and/or cell types indicates that the exons should be assigned to a single gene.

In an ninth aspect of the invention, there is provided a nucleic acid sequence as set out in any of SEQ

25 ID NOs: 1 - 25,129 wherein said sequence encodes a peptide.

In a tenth aspect of the invention, there is provided a peptide encoded by a sequence comprising a sequence as set out in any of SEQ ID NOs: 12,674 - 25,129, or a complementary sequence or coding portion thereof.

In a preferred embodiment, a peptide may be encoded by a sequence comprising a sequence set out in any of SEQ ID NOS.: 1 -12,673.

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In a further aspect, the invention provides
peptides comprising an amino acid sequence translated from
the DNA fragments, said amino acid sequences comprising SEQ

ID NOS.: 25,130 - 37,156.

Accordingly in a eleventh aspect of the invention there is provided a peptide comprising a sequence as set out in any of SEQ ID NOs: 25,130 - 37,156, or fragment thereof.

In another aspect, the invention provides means for displaying annotated sequence, and in particular, for displaying sequence annotated according to the methods and apparatus of the present invention. Further, such display can be used as a preferred graphical user interface for electronic search, query, and analysis of such annotated sequence.

15 Detailed Description of the Invention

Definitions

As used herein, the term "microarray" and phrase "nucleic acid microarray" refer to a substrate-bound collection of plural nucleic acids, hybridization to each of the plurality of bound nucleic acids being separately detectable. The substrate can be solid or porous, planar or non-planar, unitary or distributed.

As so defined, the term "microarray" and phrase

"nucleic acid microarray" include all the devices so called
in Schena (ed.), DNA Microarrays: A Practical Approach
(Practical Approach Series), Oxford University Press (1999)
(ISBN: 0199637768); Nature Genet. 21(1)(suppl):1 - 60
(1999); and Schena (ed.), Microarray Biochip: Tools and

Technology, Eaton Publishing Company/BioTechniques Books
Division (2000) (ISBN: 1881299376). As so defined, the
term "microarray" and phrase "nucleic acid microarray"
further include substrate-bound collections of plural
nucleic acids in which the nucleic acids are distributably
disposed on a plurality of beads, rather than on a unitary

planar substrate, as is described, inter alia, in Brenner et al., Proc. Natl. Acad. Sci. USA 97(4):166501670 (2000); in such case, the term "microarray" and phrase "nucleic acid microarray" refer to the plurality of beads in aggregate.

As used herein with respect to a nucleic acid microarray, the term "probe" refers to the nucleic acid that is, or is intended to be, bound to the substrate; in such context, the term "target" thus refers to nucleic acid intended to be bound thereto by Watson-Crick complementarity. As used herein with respect to solution phase hybridization, the term "probe" refers to the nucleic acid of known sequence that is detectably labeled.

As used herein, the expression "probe comprising SEQ ID NO.", and variants thereof, intends a nucleic acid probe, at least a portion of which probe has either (i) the sequence directly as given in the referenced SEQ ID NO., or (ii) a sequence complementary to the sequence as given in the referenced SEQ ID NO., the choice as between sequence directly as given and complement thereof dictated by the requirement that the probe hybridize to mRNA.

As used herein, the term "open reading frame" and the equivalent acronym "ORF" refer to that portion of an exon that can be translated in its entirety into a sequence of contiguous amino acids i.e. a nucleic acid sequence that, in at least one reading frame, does not possess stop codons; the term does not require that the ORF encode the entirety of a natural protein.

As used herein, the term "amplicon" refers to a 30 PCR product amplified from human genomic DNA, containing the predicted exon.

As used herein the term "exon" refers to the consensus prediction of the various exon and gene predicting algorithms i.e. a nucleic acid sequence bioinformatically predicted to encode a portion of a

natural protein.

As used herein, the term "peptide" refers to a sequence of amino acids. The sequences referred to as PEPTIDE SEQ ID NOS.: are the predicted peptide sequences that would be translated from one of the exons, or a portion thereof set out in exon SEQ ID NOS.:. The codons encoding the peptide are wholly contained within the exon.

As used herein, a "portions" of a defined nucleotide sequence or sequences can be and, preferably, are fragments unique to that sequence or to one or a combination of those sequences. A fragment unique to a nucleic acid molecule is one that is a signature for the larger nucleic acid molecule.

As used herein, the phrase "expression of a probe" and its linguistic variants means that the ORF present within the probe, or its complement, is present within a target mRNA.

As used herein, "stringent conditions" refers to parameters well known to those skilled in the art. When a nucleic acid molecule is said to be hybridisable to another of a given sequence under "stringent conditions" it is meant that it is homologous to the given sequence.

As used herein, the phrase "specific binding pair" intends a pair of molecules that bind to one another with high specificity. Binding pairs are said to exhibit specific binding when they exhibit avidity of at least 10⁷, preferably at least 10⁸, more preferably at least 10⁹ liters/mole. Nonlimiting examples of specific binding pairs are: antibody and antigen; biotin and avidin; and biotin and streptavidin.

As used herein with respect to the visual display of annotated genomic sequence, the term "rectangle" means any geometric shape that has at least a first and a second border, wherein the first and second borders each are capable of mapping uniquely to a point of another visual

object of the display.

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As used herein, a "Mondrian" means a visual display in which a single genomic sequence is annotated with predicted and experimentally confirmed functional information.

Brief Description of the Drawings

The present invention is further illustrated with reference to the following non-limiting figures and examples in which:

functional regions from genomic sequence, confirming the

functional activity of such regions experimentally, and
associating and displaying the data so obtained in
meaningful and useful relationship to the original sequence
data;

FIG. 2 further elaborates that portion of the 20 process schematized in FIG. 1 for predicting functional regions from genomic sequence;

FIG. 3 illustrates a Mondrian visual display;

FIG. 4 presents a Mondrian showing a hypothetical annotated genomic sequence;

FIG. 5 is a histogram showing the distribution of ORF length and PCR products as obtained, with ORF length shown in black and PCR product length shown in dotted lines;

FIG. 6 is a histogram showing the distribution,
30 among exons predicted according to the methods described,
of expression as measured using simultaneous two color
hybridization to a genome-derived single exon microarray.
The graph shows the number of sequence-verified products
that were either not expressed ("0"), expressed in one or
35 more but not all tested tissues ("1" - "9"), or expressed

in all tissues tested ("10");

FIG. 7 is a pictorial representation of the expression of verified sequences that showed expression with signal intensity greater than 3 in at least one

5 tissue, with: FIG. 7A showing the expression as measured by microarray hybridization in each of the 10 measured tissues, and the expression as measured "bioinformatically" by query of EST, NR and SwissProt databases; with FIG. 7B showing the legend for display of physical expression

10 (ratio) in FIG. 7A; and with FIG. 7C showing the legend for scoring EST hits as depicted in FIG. 7A;

FIG. 8 shows a comparison of normalized CY3 signal intensity for arrayed sequences that were identical to sequences in existing EST, NR and SwissProt databases or that were dissimilar (unknown), where black denotes the signal intensity for all sequence-verified products with a BLAST Expect ("E") value of greater than 1e-30 (1 x 10⁻³⁰) ("unknown") and a dotted line denotes sequence-verified spots with a BLAST expect ("E") value of less than 1e-30 (1 x 10⁻³⁰) ("known");

FIG. 9 presents a Mondrian of BAC AC008172 (bases 25,000 to 130,000), containing the carbamyl phosphate synthetase gene (AF154830.1); and

FIG. 10 is a Mondrian of BAC A049839.

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Methods and Apparatus for Predicting, Confirming,
Annotating, and Displaying Functional Regions From Genomic
Sequence Data

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FIG. 1 is a flow chart illustrating in broad outline a process for predicting functional regions from genomic sequence, confirming and characterizing the functional activity of such regions experimentally, and then associating and displaying the information so obtained

PCT/US01/00669 WO 01/57277

in meaningful and useful relationship to the original sequence data.

The initial input into process 10 of the present invention is drawn from one or more databases 100 5 containing genomic sequence data. Because genomic sequence is usually obtained from subgenomic fragments, the sequence data typically will be stored in a series of records corresponding to these subgenomic sequenced fragments. Some fragments will have been catenated to form larger 10 contiguous sequences ("contigs"); others will not. finite percentage of sequence data in the database will typically be erroneous, consisting inter alia of vector sequence, sequence created from aberrant cloning events, sequence of artificial polylinkers, and sequence that was 15 erroneously read.

Each sequence record in database 100 will minimally contain as annotation a unique sequence identifier (accession number), and will typically be annotated further to identify the date of accession, 20 species of origin, and depositor. Because database 100 can contain nongenomic sequence, each sequence will typically be annotated further to permit query for genomic sequence. Chromosomal origin, optionally with map location, can also be present. Data can be, and over time increasingly will 25 be, further annotated with additional information, in part through use of the present invention, as described below. Annotation can be present within the data records, in information external to database 100 and linked to the records thereto, or through a combination of the two.

Databases useful as genomic sequence database 100 in the present invention include GenBank, and particularly include several divisions thereof, including the htgs(draft), NT (nucleotide, command line), and NR (nonredundant) divisions. GenBank is produced by the 35 National Institutes of Health and is maintained by the

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PCT/US01/00669 WO 01/57277

National Center for Biotechnology Information (NCBI). Databases of genomic sequence from species other than human, such as mouse, rat, Arabidopsis, C. elegans, C. brigsii, Drosophila, zebra fish, and other higher 5 eukaryotic organisms will also prove useful as genomic sequence database 100.

Genomic sequence obtained by query of genomic sequence database 100 is then input into one or more processes 200 for identification of regions therein that 10 are predicted to have a biological function as specified by the user. Such functions include, but are not limited to, encoding protein, regulating transcription, regulating message transport after transcription into mRNA, regulating message splicing after transcription into mRNA, of 15 regulating message degradation after transcription into mRNA, and the like. Other functions include directing somatic recombination events, contributing to chromosomal stability or movement, contributing to allelic exclusion or X chromosome inactivation, and the like.

The particular genomic sequence to be input into process 200 will depend upon the function for which relevant sequence is to be identified as well as upon the approach chosen for such identification. Process step 200 can be iterated to identify different functions within a 25 given genomic region. In such case, the input often will be different for the several iterations.

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Sequences predicted to have the requisite function by process 200 are then input into process 300, where a subset of the input sequences suitable for 30 experimental confirmation is identified. Experimental confirmation can involve physical and/or bioinformatic assay. Where the subsequent experimental assay is bioinformatic, rather than physical, there are fewer constraints on the sequences that can be tested, and in 35 this latter case therefore process 300 can output the

entirety of the input sequence.

The subset of sequences output from process 300 is then used in process 400 for experimental verification and characterization of the function predicted in process 200, which experimental verification can, and often will, include both physical and bioinformatic assay.

Process 500 annotates the sequence data with the functional information obtained in the physical and/or bioinformatic assays of process 400. Such annotation can be done using any technique that usefully relates the functional information to the sequence, as, for example, by incorporating the functional data into the sequence data record itself, by linking records in a hierarchical or relational database, by linking to external databases, by a combination thereof, or by other means well known within the database arts. The data can even be submitted for incorporation into databases maintained by others, such as GenBank, which is maintained by NCBI.

As further noted in FIG. 1, additional annotation 20 can be input into process 500 from external sources 600.

The annotated data is then displayed in process 800, either before, concomitantly with, or after optional storage 700 on nontransient media, such as magnetic disk, optical disc, magnetooptical disk, flash memory, or the like.

from process 400 can be used in each preceding step of process 10: e.g., facilitating identification of functional sequences in process 200, facilitating identification of an experimentally suitable subset thereof in process 300, and facilitating creation of physical and/or informational substrates for, and performance of subsequent assay, of functional sequences in process 400.

Information from each step can be passed directly to the succeeding process, or stored in permanent or

interim form prior to passage to the succeeding process.

Often, data will be stored after each, or at least a plurality, of such process steps. Any or all process steps can be automated.

FIG. 2 further elaborates the prediction of functional sequence within genomic sequence according to process 200.

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Genomic sequence database 100 is first queried 20 for genomic sequence.

The sequence required to be returned by query 20 will depend, in the first instance, upon the function to be identified.

For example, genomic sequences that function to encode protein can be identified inter alia using gene
prediction approaches, comparative sequence analysis approaches, or combinations of the two. In gene prediction analysis, sequence from one genome is input into process 200 where at least one, preferably a plurality, of algorithmic methods are applied to identify putative coding regions. In comparative sequence analysis, by contrast, corresponding, e.g., syntenic, sequence from a plurality of sources, typically a plurality of species, is input into process 200, where at least one, possibly a plurality, of algorithmic methods are applied to compare the sequences and identify regions of least variability.

The exact content of query 20 will also depend upon the database queried. For example, if the database contains both genomic and nongenomic sequence, perhaps derived from multiple species, and the function to be determined is protein coding regions in human genomic sequence, the query will accordingly require that the sequence returned be genomic and derived from humans.

Query 20 can also incorporate criteria that compel return of sequence that meets operative requirements of the subsequent analytical method. Alternatively, or in

addition, such operative criteria can be enforced in subsequent preprocess step 24.

For example, if the function sought to be identified is protein coding, query 20 can incorporate 5 criteria that return from genomic sequence database 100 only those sequences present within contigs sufficiently long as to have obviated substantial fragmentation of any given exon among a plurality of separate sequence fragments.

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Such criteria can, for example, consist of a required minimal individual genomic sequence fragment length, such as 10 kb, more typically 20 kb, 30 kb, 40kb, and preferably 50 kb or more, as well as an optional further or alternative requirement that sequence from any 15 given clone, such as a bacterial artificial chromosome ("BAC"), be presented in no more than a finite maximal number of fragments, such as no more than 20 separate pieces, more typically no more than 15 fragments, even more typically no more than about 10 - 12 fragments.

Results using the present invention have shown that genomic sequence from bacterial artificial chromosomes (BACs) is sufficient for gene prediction analysis according to the present invention if the sequence is at least 50 kb in length, and if additionally the sequence from any given BAC is presented in fewer than 15, and preferably fewer than 10, fragments. Accordingly, query 20 can incorporate a requirement that data accessioned from BAC sequencing be in fewer than 15, preferably fewer than 10, fragments.

An additional criterion that can be incorporated 30 into the query can be the date, or range of dates, of sequence accession. Although the process has been described above as if genomic sequence database 100 were static, it is of course understood that the genomic sequence databases need not be static, and indeed are 35 typically updated on a frequent, even hourly, basis. Thus,

as further described in Examples 1 and 2, infra, it is possible to query the database for newly added sequence, either newly added after an absolute date, or newly added relative to a prior analysis performed using the methods and apparatus of the present invention. In this way, the process herein described can incorporate a dynamic, temporal component.

One utility of such temporal limitation is to identify, from newly accessioned genomic sequence, the

10 presence of novel genes, particularly those not previously identified by EST sequencing (or other sequencing efforts that are similarly based upon gene expression). As further described in Example 1, such an approach has shown that newly accessioned human genomic sequence, when analyzed for sequences that function to encode protein, readily identifies genes that are novel over those in existing EST and other expression databases. This makes the methods of the present invention extremely powerful gene discovery tools. And as would be appreciated, such gene discovery can be performed using genomic sequence from species other than human.

If query 20 incorporates multiple criteria, such as above-described, the multiple criteria can be performed as a series of separate queries or as a single query,

25 depending in part upon the query language, the complexity of the query, and other considerations well known in the database arts.

If query 20 returns no genomic sequence meeting the query criteria, the negative result can be reported by process 22, and process 200 (and indeed, entire process 10) ended 23, as shown. Alternatively, or in addition to report and termination of the initial inquiry, a new query 20 can be generated that takes into account the initial negative result.

When query 20 returns sequence meeting the query

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criteria, the returned sequence is then passed to optional preprocessing 24, suitable and specific for the desired analytical approach and the particular analytical methods thereof to be used in process 25.

Preprocessing 24 can include processes suitable for many approaches and methods thereof, as well as processes specifically suited for the intended subsequent analysis.

Preprocessing 24 suitable for most approaches and 10 methods will include elimination of sequence irrelevant to, or that would interfere with, the subsequent analysis. Such sequence includes repetitive sequence, such as Alu repeats and LINE elements, vector sequence, artificial sequence, such as artificial polylinkers, and the like. 15 Such removal can readily be performed by identification and subsequent masking of the undesired sequence.

Identification can be effected by comparing the genomic sequence returned by query 20 with public or private databases containing known repetitive sequence, 20 vector sequence, artificial sequence, and other artifactual sequence. Such comparison can readily be done using programs well known in the art, such as CROSS_MATCH, or by proprietary sequence comparison programs the engineering of which is well within the skill in the art.

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Alternatively, or in addition, undesirable, including artifactual, sequence can be identified algorithmically without comparison to external databases and thereafter removed. For example, synthetic polylinker sequence can be identified by an algorithm that identifies 30 a significantly higher than average density of known restriction sites. As another example, vector sequence can be identified by algorithms that identify nucleotide or codon usage at variance with that of the bulk of the genomic sequence.

Once identified, undesired sequence can be

removed. Removal can usefully be done by masking the undesired sequence as, for example, by converting the specific nucleotide references to one that is unrecognized by the subsequent bioinformatic algorithms, such as "X".

5 Alternatively, but at present less preferred, the undesired sequence can be excised from the returned genomic sequence, leaving gaps.

Preprocessing 24 can further include selection from among duplicative sequences of that one sequence of highest quality. Higher quality can be measured as a lower percentage of, fewest number of, or least densely clustered occurrence of ambiguous nucleotides, defined as those nucleotides that are identified in the genomic sequence using symbols indicating ambiguity. Higher quality can also or alternatively be valued by presence in the longest contig.

Preprocessing 24 can, and often will, also include formatting of the data as specifically appropriate for passage to the analytical algorithms of process 25.

20 Such formatting can and typically will include, inter alia, addition of a unique sequence identifier, either derived from the original accession number in genomic sequence database 100, or newly applied, and can further include additional annotation. Formatting can include conversion

25 from one to another sequence listing standard, such as conversion to or from FASTA or the like, depending upon the input expected by the subsequent process.

Preprocessing, which can be optional depending upon the function desired to be identified and the informational requirements of the methods for effecting such identification, is followed by sequence processing 25, where sequences with the desired function are identified within the genomic sequence.

As mentioned above, such functions can include, some are not limited to, encoding protein, regulating

transcription, regulating message transport after transcription into mRNA, regulating message splicing after transcription, of regulating message degradation, and the like. Other functions include directing somatic 5 recombination events, contributing to chromosomal stability or movement, contributing to allelic exclusion or X chromosome inactivation, or the like.

The methods of the present invention are particularly useful for gene discovery, that is, for 10 identifying, from genomic sequence, regions that function to encode genes, and in a particularly useful embodiment, for identifying regions that function to encode genes not hitherto identified by expression-based or directed cloning and sequencing. In conjunction with verification using the 15 novel single exon microarrays of the present invention, as further described below, the methods herein described become powerful gene discovery tools.

Accordingly, in a preferred embodiment of the present invention, process 25 is used to identify putative 20 coding regions. Two preferred approaches in process 25 for identifying sequence that encodes putative genes are gene prediction and comparative sequence analysis.

Gene prediction can be performed using any of a number of algorithmic methods, embodied in one or more 25 software programs, that identify open reading frames (ORFs) using a variety of heuristics, such as GRAIL, DICTION, and GENEFINDER. Comparative sequence analysis similarly can be performed using any of a variety of known programs that identify regions with lower sequence variability.

As further described in Example 1, below, gene finding software programs yield a range of results. For the newly accessioned human genomic sequence input in Example 1, for example, GRAIL identified the greatest percentage of genomic sequence as putative coding region, 35 2% of the data analyzed; GENEFINDER was second, calling 1%;

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and DICTION yielded the least putative coding region, with 0.8% of genomic sequence called as coding region.

Increased reliability can be obtained when consensus is required among several such methods. Although discussed herein particularly with respect to exon calling, consensus among methods will in general increase reliability of predicting other functions as well.

Thus, as indicated by query 26, sequence processing 25, optionally with preprocessing 24, can be repeated with a different method, with consensus among such iterations determined and reported in process 27.

Process 27 compares the several outputs for a given input genomic sequence and identifies consensus among the separately reported results. The consensus itself, as well as the sequence meeting that consensus, is then stored in process 29a, displayed in process 29b, and/or output to process 300 for subsequent identification of a subset thereof suitable for assay.

and reported by process 27. For example, as further described in Example 1, infra, process 27 can report consensus as between all specific pairs of methods of gene prediction, as consensus among any one or more of the pairs of methods of gene prediction, or as among all of the gene prediction algorithms used. Thus, in Example 1, process 27 reported that GRAIL and GENEFINDER programs agreed on 0.7% of genomic sequence, that GRAIL and DICTION agreed on 0.5% of genomic sequence, and that the three programs together agreed on 0.25% of the data analyzed. Put another way, 0.25% of the genomic sequence was identified by all three of the programs as containing putative coding region.

Furthermore, consensus can be required among different approaches to identifying a chosen function.

For example, if the function desired to be identified is coding of protein sequence, and a first used

approach to exon calling is gene prediction, the process can be repeated on the same input sequence, or subset thereof, with another approach, such as comparative sequence analysis. In such a case, where comparative sequence analysis follows gene prediction, the comparison can be performed not only on genomic nucleic acid sequence, but additionally or alternatively can be performed on the predicted amino acid sequence translated from the ORFs prior identified by the gene prediction approach.

Although shown as an iterative process, the multiple analyses required to achieve consensus can be done in series, in parallel, or some combination thereof.

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Predicted functional sequence, optionally representing a consensus among a plurality of methods and approaches for determination thereof, is passed to process 300 for identification of a subset thereof for functional assay.

In the preferred embodiment of the methods of the present invention, wherein the function sought to be

20 identified is protein coding, process 300 is used to identify a subset thereof suitable for experimental verification by physical and/or bioinformatic approaches.

For example, putative ORFs identified in process 200 can be classified, or binned, bioinformatically into putative genes. This binning can be based inter alia upon consideration of the average number of exons/gene in the species chosen for analysis, upon density of exons that have been called on the genomic sequence, and other empirical rules. Thereafter, one or more among the genespecific ORFs can be chosen for subsequent use in gene expression assay.

Where such subsequent gene expression assay uses amplified nucleic acid, considerations such as desired amplicon length, primer synthesis requirements, putative exon length, sequence GC content, existence of possible

secondary structure, and the like can be used to identify and select those ORFs that appear most likely successfully to amplify. Where subsequent gene expression assay relies upon nucleic acid hybridization, whether or not using amplified product, further considerations involving hybridization stringency can be applied to identify that subset of sequences that will most readily permit sequence-specific discrimination at a chosen hybridization and wash stringency. One particular such consideration is avoidance of putative exons that span repetitive sequence; such sequence can hybridize spuriously to nonspecific message, reducing specific signal in the hybridization.

For bioinformatic assay, there are fewer constraints on the sequences that can be tested experimentally, and in this latter case therefore process 300 can output the entirety of the input sequence.

The subset of sequences identified by process 300 as suitable for use in assay is then used in process 400 to create the physical and/or informational substrate for experimental verification of the predictions made in process 200, and thereafter to assay those substrates.

As mentioned, the methods of the present invention are particularly useful for identifying potential coding regions within genomic sequence. In a preferred embodiment of process 400, therefore, the expression of the sequences predicted to encode protein is verified. The combination of the predictive and experimental methods provides a powerful gene discovery engine.

Thus, in another aspect, the present invention

30 provides methods and apparatus for verifying the expression
of putative genes identified within genomic sequence. In
particular, the invention provides a novel method of
verifying gene expression in which expression of predicted
ORFs is measured and confirmed using a novel type of

35 nucleic acid microarray, the genome-derived single exon

nucleic acid microarrays of the present invention.

Putative ORFs as predicted by a consensus of gene calling, particularly gene prediction, algorithms in process 200, and as further identified as suitable by process 300, are amplified from genomic DNA using the polymerase chain reaction (PCR). Although PCR is conveniently used, other amplification approaches can also be used.

Amplification schemes can be designed to capture
the entirety of each predicted ORF in an amplicon with
minimal additional (that is, intronic or intergenic)
sequence. Because ORFs predicted from human genomic
sequence using the methods of the present invention differ
in length, such an approach results in amplicons of varying
length.

However, most predicted ORFs are shorter than 500 bp in length, and although amplicons of at least about 100 or 200 base pairs can be immobilized as probes on nucleic acid microarrays, early experimental results using the

20 methods of the present invention have suggested that longer amplicons, at least about 400 or 500 base pairs, are more effective. Furthermore, certain advantages derive from application to the microarray of amplicons of defined size.

Therefore, amplification schemes can

25 alternatively, and preferably, be designed to amplify regions of defined size, preferably at least about 300, 400 or 500 bp, centered about each predicted ORF. Such an approach results in a population of amplicons of limited size diversity, but that typically contain intronic and/or intergenic nucleic acid in addition to putative ORF.

Conversely, somewhat fewer than 10% of ORFs predicted from human genomic sequence according to the methods of the present invention exceed 500 bp in length.

Portions of such extended ORFs, preferably at least about 300,400 or 500 bp in length, can be amplified. However, it

has been discovered that the percentage success at amplifying pieces of such ORFs is low, and that such putative exons are more effectively amplified when larger fragments, at least about 1000 or 1500 bp, and even as large as 2000 bp are amplified.

The putative ORFs selected in process 300 are thus input into one or more primer design programs, such as PRIMER3 (available online for use at http://www-genome.wi.mit.edu/cgi-bin/primer/), with a goal of amplifying at least about 500 base pairs of genomic sequence centered within or about ORFs predicted to be no more than about 500 bp, or at least about 1000 - 1500 bp of genomic sequence for ORFs predicted to exceed 500 bp in length, and the primers synthesized by standard techniques.

15 Primers with the requisite sequences can be purchased commercially or synthesized by standard techniques.

Conveniently, a first predetermined sequence can be added commonly to the ORF-specific 5' primer and a second, typically different, predetermined sequence

commonly added to each 3' ORF-unique primer. This serves to immortalize the amplicon, that is, serves to permit further amplification of any amplicon using a single set of primers complementary respectively to the common 5' and common 3' sequence elements. The presence of these

"universal" priming sequences further facilitates later sequence verification, providing a sequence common to all amplicons at which to prime sequencing reactions. The common 5' and 3' sequences further serve to add a cloning site should any of the ORFs warrant further study.

Such predetermined sequence is usefully at least about 10, 12 or 15 nt in length, and usually does not exceed about 25 nt in length. The "universal" priming sequences used in the examples presented *infra* were each 16 nt long.

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The genomic DNA to be used as substrate for

amplification will come from the eukaryotic species from which the genomic sequence data had originally been obtained, or a closely related species, and can conveniently be prepared by well known techniques from somatic or germline tissue or cultured cells of the organism. See, e.g., Short Protocols in Molecular Biology: A Compendium of Methods from Current Protocols in Molecular Biology, Ausubel et al. (eds.), 4th edition (April 1999), John Wiley & Sons (ISBN: 047132938X) and Maniatis et al., Molecular Cloning: A Laboratory Manual, 2nd edition (December 1989), Cold Spring Harbor Laboratory Press (ISBN: 0879693096). Many such prepared genomic DNAs are available commercially, with the human genomic DNAs additionally having certification of donor informed

Although the intronic and intergenic material flanking putative coding regions in the amplicons could potentially interfere with hybridizations during microarray experiments, we have found, surprisingly, that differential expression ratios are not significantly affected. Rather, the predominant effect of exon size is to alter the absolute signal intensity, rather than its ratio. Equally surprising, the art had suggested that single exon probes would not provide sufficient signal intensity for high stringency hybridization analyses; we find that such probes not only provide adequate signal, but have substantial advantages, as herein described.

After partial purification, as by size exclusion spin column, with or without confirmation as to amplicon quality as by gel electrophoresis, each amplicon (single exon probe) is disposed in an array upon a support substrate.

Methods for creating microarrays by deposition and fixation of nucleic acids onto support substrates are well known in the art (Reviewed by Schena et al., see

above).

Typically, the support substrate will be glass, although other materials, such as amorphous or crystalline silicon or plastics. Such plastics include

5 polymethylacrylic, polyethylene, polypropylene, polyacrylate, polymethylmethacrylate, polyvinylchloride, polytetrafluoroethylene, polystyrene, polycarbonate, polyacetal, polysulfone, celluloseacetate, cellulosenitrate, nitrocellulose, or mixtures thereof, can also be used. Typically, the support will be rectangular, although other shapes, particularly circular disks and even spheres, present certain advantages. Particularly advantageous alternatives to glass slides as support substrates for array of nucleic acids are optical discs, as described in WO 98/12559.

The amplified nucleic acids can be attached covalently to a surface of the support substrate or, more typically, applied to a derivatized surface in a chaotropic agent that facilitates denaturation and adherence by presumed noncovalent interactions, or some combination thereof.

Robotic spotting devices useful for arraying nucleic acids on support substrates can be constructed using public domain specifications (The MGuide, version 2.0, http://cmgm.stanford.edu/pbrown/mguide/index.html), or can conveniently be purchased from commercial sources (MicroArray GenII Spotter and MicroArray GenIII Spotter, Molecular Dynamics, Inc., Sunnyvale, CA). Spotting can also be effected by printing methods, including those using ink jet technology.

As is well known in the art, microarrays typically also contain immobilized control nucleic acids. For controls useful in providing measurements of background signal for the genome-derived single exon microarrays of the present invention, a plurality of *E. coli* genes can

readily be used. As further described in Example 1, 16 or 32 E. coli genes suffice to provide a robust measure of background noise in such microarrays.

As is well known in the art, the amplified 5 product disposed in arrays on a support substrate to create a nucleic acid microarray can consist entirely of natural nucleotides linked by phosphodiester bonds, or alternatively can include either nonnative nucleotides, alternative internucleotide linkages, or both, so long as 10 complementary binding can be obtained in the hybridization. If enzymatic amplification is used to produce the immobilized probes, the amplifying enzyme will impose certain further constraints upon the types of nucleic acid analogs that can be generated.

Although particularly described herein as using high density microarrays constructed on planar substrates, the methods of the present invention for confirming the expression of ORFs predicted from genomic sequence can use any of the known types of microarrays, as herein defined, 20 including lower density planar arrays, and microarrays on nonplanar, nonunitary, distributed substrates.

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For example, gene expression can be confirmed using hybridization to lower density arrays, such as those constructed on membranes, such as nitrocellulose, nylon, 25 and positively-charged derivatized nylon membranes. Further, gene expression can also be confirmed using nonplanar, bead-based microarrays such as are described in Brenner et al., Proc. Natl. Acad. Sci. USA 97(4):166501670 (2000); U.S. Patent No. 6,057,107; and U.S. Patent No. 30 5,736,330. In theory, a packed collection of such beads provides in aggregate a higher density of nucleic acid probe than can be achieved with spotting or lithography techniques on a single planar substrate.

Planar microarrays on solid substrates, however, 35 provide certain useful advantages, including high

throughput and compatibility with existing readers. For example, each standard microscope slide can include at least 1000, typically at least 2000, preferably 5000 and upto 10,000 - 50,000 or more nucleic acid probes of discrete sequence. The number of sequences deposited will depend on their required application.

Each putative gene can be represented in the array by a single predicted ORF. Alternatively, genes can be represented by more than one predicted ORF. For purposes of measuring differential splicing, more than one predicted ORF will be provided for a putative gene. And as is well known in the art, each probe of defined sequence, representing a single predicted ORF, can be deposited in a plurality of locations on a single microarray to provide redundancy of signal.

The genome-derived single exon microarrays described above differ in several fundamental and advantageous ways from microarrays presently used in the gene expression art, including (1) those created by deposition of mRNA-derived nucleic acids, (2) those created by in situ synthesis of oligonucleotide probes, and (3) those constructed from yeast genomic DNA.

Most nucleic acid microarrays that are in use for study of eukaryotic gene expression have as immobilized probes nucleic acids that are derived — either directly or indirectly — from expressed message. As discussed above, it is common, for example, for such microarrays to be derived from cDNA/EST libraries, either from those previously described in the literature, see Lennon et al., or from the de novo construction of "problem specific" libraries targeted at a particular biological question, R.S. Thomas et al., Cancer Res. (in press). Such microarrays are herein collectively denominated "EST microarrays".

Such EST microarrays by definition can measure

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expression only of those genes found in EST libraries, shown herein to represent only a fraction of expressed genes. Furthermore, such libraries - and thus microarrays based thereupon - are biased by the tissue or cell type of 5 message origin, by the expression levels of the respective genes within the tissues, and by the ability of the message successfully to have been reverse-transcribed and cloned.

Thus, as further discussed in Example 1, the methods of the present invention enable sequences that do 10 not appear in EST or other expression databases to be determined - subsequently arrayed for expression measurements could not, therefore, have been represented as probes on an EST microarray. And as further demonstrated in the examples, infra, the remaining population of genes 15 identified from genomic sequence by the methods of the present invention - that is, the one third of sequences that had previously been accessioned in EST or other expression databases - are biased toward genes with higher expression levels.

Representation of a message in an EST and/or cDNA library depends upon the successful reverse transcription, optionally but typically with subsequent successful cloning, of the message. This introduces substantial bias into the population of probes available for arraying in EST 25 microarrays.

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In contrast, neither reverse transcription nor cloning is required to produce the probes arrayed on the genome-derived single exon microarrays of the present invention. And although the ultimate deposition of a probe 30 on the genome-derived single exon microarray of the present invention depends upon a successful amplification from genomic material, a priori knowledge of the sequence of the desired amplicon affords greater opportunity to recover any given probe sequence recalcitrant to amplification than is 35 afforded by the requirement for successful reverse

transcription and cloning of unknown message in EST approaches.

Thus, the genome-derived single exon microarrays of the present invention present a far greater diversity of 5 probes for measuring gene expression, with far less bias, than do EST microarrays presently used in the art.

As a further consequence of their ultimate origin from expressed message, the probes in EST microarrays often contain poly-A (or complementary poly-T) stretches derived 10 from the poly-A tail of mature mRNA. These homopolymeric stretches contribute to cross-hybridization, that is, to a spurious signal occasioned by hybridization to the homopolymeric tail of a labeled cDNA that lacks sequence homology to the gene-specific portion of the probe.

In contrast, the probes arrayed in the genomederived single exon microarrays of the present invention lack homopolymeric stretches derived from message polyadenylation, and thus can provide more specific signal. Typically, at least about 50, 60 or 75% of the probes on 20 the genome-derived single exon microarrays of the present invention lack homopolymeric regions consisting of A or T, where a homopolymeric region is defined for purposes herein as stretches of 25 or more, typically 30 or more, identical nucleotides.

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A further distinction, which also affects the specificity of hybridization, is occasioned by the typical derivation of EST microarray probes from cloned material. Because much of the probe material disposed as probes on EST microarrays is excised or amplified from plasmid, 30 phage, or phagemid vectors, EST microarrays typically include a fair amount of vector sequence, more so when the probes are amplified, rather than excised, from the vector.

In contrast, the vast majority of probes in the genome-derived single exon microarrays of the present invention contain no prokaryotic or bacteriophage vector

sequence, having been amplified directly or indirectly from genomic DNA. Typically, therefore, at least about 50, 60, 70 or 80% or more of individual exon-including probes disposed on a genome-derived single exon microarray of the present invention lack vector sequence, and particularly lack sequences drawn from plasmids and bacteriophage. Preferably, at least about 85, 90 or more than 90% of exon-including probes in the genome-derived single exon microarray of the present invention lack vector sequence.

With attention to removal of vector sequences through preprocessing 24, percentages of vector-free exon-including

preprocessing 24, percentages of vector-free exon-including probes can be as high as 95 - 99%. The substantial absence of vector sequence from the genome-derived single exon microarrays of the present invention results in greater specificity during hybridization, since spurious cross-hybridization to a probe vector sequence is reduced.

As a further consequence of excision or amplification of probes from vectors in construction of EST microarrays, the probes arrayed thereon often contain artificial sequence, derived from vector polylinker multiple cloning sites, at both 5' and 3' ends. The probes disposed upon the genome-derived single exon microarrays need have no such artificial sequence appended thereto.

As mentioned above, however, the ORF-specific

primers used to amplify putative ORFs can include
artificial sequences, typically 5' to the ORF-specific
primer sequence, useful for "universal" (that is,
independent of ORF sequence) priming of subsequent
amplification or sequencing reactions. When such

"universal" 5' and/or 3' priming sequences are appended to
the amplification primers, the probes disposed upon the
genome-derived single exon microarray will include
artificial sequence similar to that found in EST
microarrays. However, the genome-derived single exon
microarray of the present invention can be made without

such sequences, and if so constructed, presents an even smaller amount of nonspecific sequence that would contribute to nonspecific hybridization.

Yet another consequence of typical use of cloned 5 material as probes in EST microarrays is that such microarrays contain probes that result from cloning artifacts, such as chimeric molecules containing coding region of two separate genes. Derived from genomic material, typically not thereafter cloned, the probes of 10 the genome-derived single exon microarrays of the present invention lack such cloning artifacts, and thus provide greater specificity of signal in gene expression measurements.

A further consequence of the cloned origin of 15 probes on many EST microarrays is that the individual probes often have disparate sizes, which can cause the optimal hybridization stringency to vary among probes on a single microarray. In contrast, as discussed above, the probes arrayed on the genome-derived single exon 20 microarrays of the present invention can readily be designed to have a narrow distribution in sizes, with the range of probe sizes no greater than about 10% of the average size, typically no greater than about 5% of the average probe size.

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Because of their origin from fully- or partiallyspliced message, probes disposed upon EST arrays will often include multiple exons. The percentage of such exonspanning probes in an EST microarray can be calculated, on average, based upon the predicted number of exons/gene for 30 the given species and the average length of the immobilized probes. For human genes, the near-complete sequence of human chromosome 22, Dunham et al., Nature 402(6761):489-95 (1999), predicts that human genes average 5.5 exons/gene. Even with probes of 200 - 500 bp, the vast majority of 35 human EST microarray probes include more than one exon.

In contrast, by virtue of their origin from algorithmically identified ORFs in genomic sequence, the probes in the genome-derived single exon microarrays of the present invention can consist of individual exons. Thus, in contrast to EST microarrays, at least about 50, 60, 70, 75, 80, 85, 95 or 99% of probes deposited in the genomederived microarray of the present invention consist of, or include, no more than one predicted ORF.

This provides the ability, not readily achieved using EST microarrays, to use the genome-derived single exon microarrays of the present invention to measure tissue-specific expression of individual exons, which in turn allows differential splicing events to be detected and characterized, and in particular, allows the correlation of differential splicing to tissue-specific expression patterns.

Furthermore, the exons that are represented in EST microarrays are often biased toward the 3' or 5' end of their respective genes, since sequencing strategies used

20 for EST identification are so biased. In contrast, no such 3' or 5' bias necessarily inheres in the selection of exons for disposition on the genome-derived single exon microarrays of the present invention.

Conversely, the probes provided on the genomederived single exon microarrays of the present invention
typically, but need not necessarily, include intronic
and/or intergenic sequence that is absent from EST
microarrays, which are derived from mature mRNA.
Typically, at least about 50, 60, 70, 80 or 90% of the
exon-including probes on the genome-derived single exon
microarrays of the present invention include sequence drawn
from noncoding regions. As discussed above, the additional
presence of noncoding region does not significantly
interfere with measurement of gene expression, and provides
the additional opportunity to assay prespliced RNA, and

thus measure such phenomena such as nuclear export control.

The genome-derived single exon microarrays of the present invention are also quite different from in situ synthesis microarrays, where probe size is severely constrained by inadequacies in the photolithographic synthesis process.

Typically, probes arrayed on in situ synthesis microarrays are limited to a maximum of about 25 bp. As a well known consequence, hybridization to such chips must be performed at low stringency. In order, therefore, to achieve unambiguous sequence-specific hybridization results, the in situ synthesis microarray requires substantial redundancy, with concomitant programmed arraying for each probe of probe analogues with altered (i.e., mismatched) sequence.

In contrast, the longer probe length of the genome-derived single exon microarrays of the present invention allows much higher stringency hybridization and wash. Typically, therefore, exon-including probes on the genome-derived single exon microarrays of the present invention average at least about 100, 200, 300, 400 or 500 bp in length. By obviating the need for substantial probe redundancy, this approach permits a higher density of probes for discrete exons or genes to be arrayed on the microarrays of the present invention than can be achieved for in situ synthesis microarrays.

A further distinction is that the probes in in situ synthesis microarrays typically are covalently linked to the substrate surface. In contrast, the probes disposed on the genome-derived microarray of the present invention typically are; but need not necessarily be, bound noncovalently to the substrate.

Furthermore, the short probe size on *in situ* microarrays causes large percentage differences in the melting temperature of probes hybridized to their

PCT/US01/00669 WO 01/57277

complementary target sequence, and thus causes large percentage differences in the theoretically optimum stringency across the array as a whole.

In contrast, the larger probe size in the 5 microarrays of the present invention create lower percentage differences in melting temperature across the range of arrayed probes.

A further significant advantage of the microarrays of the present invention over in situ 10 synthesized arrays is that the quality of each individual probe can be confirmed before deposition. In contrast, the quality of probes cannot be assessed on a probe-by-probe basis for the in situ synthesized microarrays presently being used.

The genome-derived single exon microarrays of the 15 present invention are also distinguished over, and present substantial benefits over, the genome-derived microarrays from lower eukaryotes such as yeast. Lashkari et al., Proc. Natl. Acad. Sci. USA 94:13057-13062 (1997).

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Only about 220 - 250 of the 6100 or so nuclear genes in Saccharomyces cerevisiae - that is, only about 4 - 5% - have standard, spliceosomal, introns, Lopez et al., Nucl. Acids Res. 28:85-86 (2000); Spingola et al., RNA 5(2):221-34 (1999). Furthermore, the entire yeast genome 25 has already been sequenced. These two facts permit the ready amplification and disposition of single-ORF amplicons on such microarray without the requirement for antecedent use of gene prediction and/or comparative sequence analyses.

Thus, a significant aspect of the present invention is the ability to identify and to confirm expression of predicted coding regions in genomic sequence drawn from eukaryotic organisms that have a higher percentage of genes having introns than do yeast such as Saccharomyces cerevisiae, particularly in genomic sequence

PCT/US01/00669 WO 01/57277

drawn from eukaryotes in which at least about 10, 20 or 50% of protein-encoding genes have introns. In preferred embodiments, the methods and apparatus of the present invention are used to identify and confirm expression of 5 novel genes from genomic sequence of eukaryotes in which the average number of introns per gene is at least about one, two or three or more.

After the physical substrate is prepared, experimental verification of predicted function is 10 performed.

In a preferred embodiment of the present invention, where the function sought to be identified in genomic sequence is protein coding, experimental verification is performed by measuring expression of the 15 putative ORFs, typically through nucleic acid hybridization experiments, and in particularly preferred embodiments, through hybridization to genome-derived single exon microarrays prepared as above- described.

Expression is conveniently measured and expressed 20 for each probe in the microarray as a ratio of the expression measured concurrently in a plurality of mRNA sources, according to techniques well known in the microarray art, Reviewed in Schena et al., and as further described in Example 2, below. The mRNA source for the 25 reference against which specific expression is measured can be drawn from a homogeneous mRNA source, such as a single cultured cell-type, or alternatively can be heterogeneous, as from a pool of mRNA derived from multiple tissues and/or cell types, as further described in Example 2, infra.

mRNA can be prepared by standard techniques, see Ausubel et al. and Maniatis et al., or purchased commercially. The mRNA is then typically reversetranscribed in the presence of labeled nucleotides: the index source (that in which expression is desired to be 35 measured) is reverse transcribed in the presence of

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nucleotides labeled with a first label, typically a fluorophore (fluorochrome; fluor; fluorescent dye); the reference source is reverse transcribed in the presence of a second label, typically a fluorophore, typically fluorometrically-distinguishable from the first label. As further described in Example 2, infra, Cy3 and Cy5 dyes

prove particularly useful in these methods. After partial purification of the index and reference targets, hybridization to the probe array is conducted according to standard techniques, typically under a coverslip.

After wash, microarrays are conveniently scanned using a commercial microarray scanning device, such as a Gen3 Scanner (Molecular Dynamics, Sunnyvale, CA). Data on expression is then passed, with or without interim storage, to process 500, where the results for each probe are related to the original sequence.

Often, hybridization of target material to the genome-derived single exon microarray will identify certain of the probes thereon as of particular interest. Thus, it is often desirable that the user be able readily to obtain sufficient quantities of an individual probe, either for subsequent arrayed deposition upon an additional support substrate, often as part of a microarray having a plurality of probes so identified, or alternatively or additionally as a solitary solid-phase or solution-phase probe, for further use.

Thus, in another aspect, the present invention provides compositions and kits for the ready production of nucleic acids identical in sequence to, or substantially identical in sequence to, probes on the genome-derived single exon microarrays of the present invention.

In this aspect, a small quantity of each probe is disposed, typically without attachment to substrate, in a spatially-addressable ordered set, typically one per well of a microtiter dish. Although a 96 well microtiter plate

can be used, greater efficiency is obtained using higher density arrays, such as are provided by microtiter plates having 384, 864, 1536, 3456, 6144, or 9600 wells, and although microtiter plates having physical depressions (wells) are conveniently used, any device that permits addressable withdrawal of reagent from fluidly-noncommunicating areas can be used.

In this aspect of the invention, therefore, a fluidly noncommunicating addressable ordered set of individual probes, corresponding to those on a genomederived single exon microarray, is provided, with each probe in sufficient quantity to permit amplification, such as by PCR. As earlier mentioned, the ORF-specific 5' primers used for genomic amplification can have a first common sequence added thereto, and the ORF-specific 3' primers used for genomic amplification can have a second, different, common sequence added thereto, thus permitting, in this preferred embodiment, the use of a single set of 5' and 3' primers to amplify any one of the probes from the amplifiable ordered set.

Each discrete amplifiable probe can also be packaged with amplification primers, solutes, buffers, etc., and can be provided in dry (e.g., lyophilized) form or wet, in the latter case typically with addition of agents that retard evaporation.

In another aspect of the present invention, a genome-derived single-exon microarray is packaged together with such an ordered set of amplifiable probes corresponding to the probes, or one or more subsets of probes, thereon. In alternative embodiments, the ordered set of amplifiable probes is packaged separately from the genome-derived single exon microarray.

In some embodiments, the microarray and/or ordered probe set are further packaged with recordable media that provide probe identification and addressing

information, and that can additionally contain annotation information, such as gene expression data. Such recordable media can be packaged with the microarray, with the ordered probe set, or with both.

If the microarray is constructed on a substrate that incorporates recordable media, such as is described in international patent application no. WO 98/12559, then separate packaging of the genome-derived single exon microarray and the bioinformatic information is not required.

The amount of amplifiable probe material should be sufficient to permit at least one amplification sufficient for subsequent hybridization assay.

Although the use of high density genome-derived microarrays on solid planar substrates is presently a preferred approach for the physical confirmation and characterization of the expression of sequences predicted to encode protein, other types of microarrays (as herein defined) can also be used.

Furthermore, as earlier mentioned, experimental verification of the function predicted from genomic sequence in process 200 can be bioinformatic, rather than, or additional to, physical verification.

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For example, where the function desired to be
identified is protein coding, the predicted ORFs can be
compared bioinformatically to sequences known or suspected
of being expressed.

Thus, the sequences output from process 300 (or process 200), can be used to query expression databases,

30 such as EST databases, SNP ("single nucleotide polymorphism") databases, known cDNA and mRNA sequences,

SAGE ("serial analysis of gene expression") databases, and more generalized sequence databases that allow query for expressed sequences. Such query can be done by any

35 sequence query algorithm, such as BLAST ("basic local")

alignment search tool"). The results of such query — including information on identical sequences and information on nonidentical sequences that have diffuse or focal regions of sequence homology to the query sequence — 5 can then be passed directly to process 500, or used to inform analyses subsequently undertaken in process 200, process 300, or process 400.

Experimental data, whether obtained by physical or bioinformatic assay in process 400, is passed to process 500 where it is usefully related to the sequence data itself, a process colloquially termed "annotation". Such annotation can be done using any technique that usefully relates the functional information to the sequence, as, for example, by incorporating the functional data into the record itself, by linking records in a hierarchical or relational database, by linking to external databases, or by a combination thereof. Such database techniques are well within the skill in the art.

The annotated sequence data can be stored
locally, uploaded to genomic sequence database 100, and/or displayed 800.

The methods and apparatus of the present invention rapidly produce functional information from genomic sequence. Coupled with the escalating pace at which sequence now accumulates, the rapid pace of sequence annotation produces a need for methods of displaying the information in meaningful ways.

FIG. 3 shows visual display 80 presenting a single genomic sequence annotated according to the present invention. Because of its nominal resemblance to artistic works of Piet Mondrian, visual display 80 is alternatively described herein as a "Mondrian".

Each of the visual elements of display 80 is aligned with respect to the genomic sequence being annotated (hereinafter, the "annotated sequence"). Given

the number of nucleotides typically represented in an annotated sequence, representation of individual nucleotides would rarely be readable in hard copy output of display 80. Typically, therefore, the annotated sequence is schematized as rectangle 89, extending from the left border of display 80 to its right border. By convention herein, the left border of rectangle 89 represents the first nucleotide of the sequence and the right border of rectangle 89 represents the last nucleotide of the sequence.

As further discussed below, however, the Mondrian visual display of annotated sequence can serve as a convenient graphical user interface for computerized representation, analysis, and query of information stored electronically. For such use, the individual nucleotides can conveniently be linked to the X axis coordinate of rectangle 89. This permits the annotated sequence at any point within rectangle 89 readily to be viewed, either automatically — for example, by time-delayed appearance of a small overlaid window upon movement of a cursor or other pointer over rectangle 89 — or through user intervention, as by clicking a mouse or other pointing device at a point in rectangle 89.

Visual display 80 is generated after user

specification of the genomic sequence to be displayed.

Such specification can consist of or include an accession number for a single clone (e.g., a single BAC accessioned into GenBank), wherein the starting and stopping nucleotides are thus absolutely identified, or

alternatively can consist of or include an anchor or fulcrum point about which a chosen range of sequence is anchored, thus providing relative endpoints for the sequence to be displayed. For example, the user can anchor such a range about a given chromosomal map location, gene

name, or even a sequence returned by query for similarity

or identity to an input query sequence. When visual display 80 is used as a graphical user interface to computerized data, additional control over the first and last displayed nucleotide will typically be dynamically selectable, as by use of standard zooming and/or selection tools.

Field 81 of visual display 80 is used to present the output from process 200, that is, to present the bioinformatic prediction of those sequences having the desired function within the genomic sequence. Functional sequences are typically indicated by at least one rectangle 83 (83a, 83b, 83c), the left and right borders of which respectively indicate, by their X-axis coordinates, the starting and ending nucleotides of the region predicted to have function.

Where a single bioinformatic method or approach identifies a plurality of regions having the desired function, a plurality of rectangles 83 is disposed horizontally in field 81. Where multiple methods and/or approaches are used to identify function, each such method and/or approach can be represented by its own series of horizontally disposed rectangles 83, each such horizontally disposed series of rectangles offset vertically from those representing the results of the other methods and approaches.

Thus, rectangles 83a in FIG. 3 represent the functional predictions of a first method of a first approach for predicting function, rectangles 83b represent the functional predictions of a second method and/or second approach for predicting that function, and rectangles 83c represent the predictions of a third method and/or approach.

Where the function desired to be identified is protein coding, field 81 is used to present the bioinformatic prediction of sequences encoding protein.

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For example, rectangles 83a can represent the results from GRAIL or GRAIL II, rectangles 83b can represent the results from GENEFINDER, and rectangles 83c can represent the results from DICTION.

Optionally, and preferably, rectangles 83 collectively representing predictions of a single method and/or approach are identically colored and/or textured, and are distinguishable from the color and/or texture used for a different method and/or approach.

Alternatively, or in addition, the color, hue, density, or texture of rectangles 83 can be used further to report a measure of the bioinformatic reliability of the prediction. For example, many gene prediction programs will report a measure of the reliability of prediction.

Thus, increasing degrees of such reliability can be indicated, e.g., by increasing density of shading. Where display 80 is used as a graphical user interface, such measures of reliability, and indeed all other results output by the program, can additionally or alternatively be made accessible through linkage from individual rectangles 83, as by time-delayed window ("tool tip" window), or by pointer (e.g., mouse)-activated link.

As earlier described, increased predictive reliability can be achieved by requiring consensus among methods and/or approaches to determining function. Thus, field 81 can include a horizontal series of rectangles 83 that indicate one or more degrees of consensus in predictions of function.

Although FIG. 3 shows three series of

horizontally disposed rectangles in field 81, display 80

can include as few as one such series of rectangles and as

many as can discriminably be displayed, depending upon the

number of methods and/or approaches used to predict a given

function.

Furthermore, field 81 can be used to show

predictions of a plurality of different functions.

However, the increased visual complexity occasioned by such display makes more useful the ability of the user to select a single function for display. When display 80 is used as a graphical user interface for computer query and analysis, such function can usefully be indicated and user-selectable, as by a series of graphical buttons or tabs (not shown in FIG. 3).

Rectangle 89 is shown in FIG. 3 as including
interposed rectangle 84. Rectangle 84 represents the
portion of annotated sequence for which predicted
functional information has been assayed physically, with
the starting and ending nucleotides of the assayed material
indicated by the X axis coordinates of the left and right
borders of rectangle 84. Rectangle 85, with optional
inclusive circles 86 (86a, 86b, and 86c) displays the
results of such physical assay.

Although a single rectangle 84 is shown in FIG.

3, physical assay is not limited to just one region of

20 annotated genomic sequence. It is expected that an
 increasing percentage of regions predicted to have function
 by process 200 will be assayed physically, and that display
 80 will accordingly, for any given genomic sequence, have
 an increasing number of rectangles 84 and 85, representing

25 an increased density of sequence annotation.

Where the function desired to be identified is protein coding, rectangle 84 identifies the sequence of the probe used to measure expression. In embodiments of the present invention where expression is measured using genome-derived single exon microarrays, rectangle 84 identifies the sequence included within the probe immobilized on the support surface of the microarray. As noted supra, such probe will often include a small amount of additional, synthetic, material incorporated during amplification and designed to permit reamplification of the

probe, which sequence is typically not shown in display 80.

Rectangle 87 is used to present the results of bioinformatic assay of the genomic sequence. For example, where the function desired to be identified is protein 5 coding, process 400 can include bioinformatic query of expression databases with the sequences predicted in process 200 to encode exons. And as earlier discussed, because bioinformatic assay presents fewer constraints than does physical assay, often the entire output of process 200 10 can be used for such assay, without further subsetting thereof by process 300. Therefore, rectangle 87 typically need not have separate indicators therein of regions submitted for bioinformatic assay; that is, rectangle 87 typically need not have regions therein analogous to 15 rectangles 84 within rectangle 89.

Rectangle 87 as shown in FIG. 3 includes smaller rectangles 880 and 88. Rectangles 880 indicate regions that returned a positive result in the bioinformatic assay, with rectangles 88 representing regions that did not return 20 such positive results. Where the function desired to be predicted and displayed is protein coding, rectangles 880 indicate regions of the predicted exons that identify sequence with significant similarity in expression databases, such as EST, SNP, SAGE databases, with 25 rectangles 88 indicating genes novel over those identified in existing expression data bases.

Rectangles 880 can further indicate, through color, shading, texture, or the like, additional information obtained from bioinformatic assay.

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For example, where the function assayed and displayed is protein coding, the degree of shading of rectangles 880 can be used to represent the degree of sequence similarity found upon query of expression databases. The number of levels of discrimination can be 35 as few as two (identity, and similarity, where similarity

has a user-selectable lower threshold). Alternatively, as many different levels of discrimination can be indicated as can visually be discriminated.

Where display 80 is used as a graphical user

interface, rectangles 880 can additionally provide links
directly to the sequences identified by the query of
expression databases, and/or statistical summaries thereof.
As with each of the precedingly-discussed uses of display
80 as a graphical user interface, it should be understood
that the information accessed via display 80 need not be
resident on the computer presenting such display, which
often will be serving as a client, with the linked
information resident on one or more remotely located
servers.

Rectangle 85 displays the results of physical assay of the sequence delimited by its left and right borders.

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Rectangle 85 can consist of a single rectangle, thus indicating a single assay, or alternatively, and increasingly typically, will consist of a series of rectangles (85a, 85b, 85c) indicating separate physical assays of the same sequence.

Where the function assayed is gene expression, and where gene expression is assayed as herein described using simultaneous two-color fluorescent detection of hybridization to genome-derived single exon microarrays, individual rectangles 85 can be colored to indicate the degree of expression relative to control. Conveniently, shades of green can be used to depict expression in the sample over control values, and shades of red used to depict expression less than control, corresponding to the spectra of the Cy3 and Cy5 dyes conventionally used for respective labeling thereof. Additional functional information can be provided in the form of circles 86 (86a, 86b, 86c), where the diameter of the circle can be used to

indicate expression intensity. As discussed *infra*, such relative expression (expression ratios) and absolute expression (signal intensity) can be expressed using normalized values.

5 Where display 80 is used as a graphical user interface, rectangle 85 can be used as a link to further information about the assay. For example, where the assay is one for gene expression, each rectangle 85 can be used to link to information about the source of the hybridized mRNA, the identity of the control, raw or processed data from the microarray scan, or the like.

gene prediction and gene expression for a hypothetical BAC, showing conventions used in the Examples presented infra.

BAC sequence ("Chip seq.") 89 is presented, with the physically assayed region thereof (corresponding to rectangle 84 in FIG. 3) shown in white. Algorithmic gene predictions are shown in field 81, with predictions by GRAIL shown, predictions by GENEFINDER, and predictions by DICTION shown. Within rectangle 87, regions of sequence that, when used to query expression databases, return identical or similar sequences ("EST hit") are shown as white rectangles (corresponding to rectangles 880 in FIG. 3), gray indicates low homology, and black indicates unknowns (where black and gray would correspond to rectangles 88 in FIG. 3).

Although FIGS. 3 and 4 show a single stretch of sequence, uninterrupted from left to right, longer sequences are usefully represented by vertical stacking of such individual Mondrians, as shown in FIGS. 9 and 10.

Single Exon Probes Useful For Measuring Gene Expression

The methods and apparatus of the present invention rapidly produce functional information from

genomic sequence. Where the function to be identified is protein coding, the methods and apparatus of the present invention rapidly identify and confirm the expression of portions of genomic sequence that function to encode 5 protein. As a direct result, the methods and apparatus of the present invention rapidly yield large numbers of single-exon nucleic acid probes, the majority from previously unknown genes, each of which is useful for measuring and/or surveying expression of a specific gene in 10 one or more tissues or cell types.

It is, therefore, another aspect of the present invention to provide genome-derived single exon nucleic acid probes useful for gene expression analysis, and particularly for gene expression analysis by microarray.

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Using the methods and genome-derived single-exon microarrays of the present invention, we have for example readily identified a large number of unique ORFs from human genomic sequence. Using single exon probes that encompass these ORFs, we have demonstrated, through microarray 20 hybridization analysis, the expression of 12,673 of these ORFs in Fetal liver.

As would immediately be appreciated by one of skill in the art, each single exon probe having demonstrable expression in Fetal liver is currently 25 available for use in measuring the level of its ORF's expression in Fetal liver.

Diseases of the liver are a significant cause of human morbidity and mortality. Increasingly, genetic factors are being found that contribute to predisposition, 30 onset, and/or aggressiveness of most, if not all, of these diseases; although causative mutations in single genes have been identified for some, these disorders are believed for the most part to have polygenic etiologies.

For example, cirrhosis is a major public health 35 problem. In the industrialized world, it is among the top

ten causes of death; among patients aged 45 to 65, it is the third leading cause of death. The high prevalence is largely the result of alcohol abuse, but other major contributors include chronic $\ensuremath{\zeta_{ij}}$ hepatitis, biliary disease and 5 iron overload. Approximately 10-15% are cryptogenic.

Cirrhosis is a broad description encompassing the common end stage of many forms of liver injury. Many patients with cirrhosis will remain asymptomatic for years, while others show generalized weakness, anorexia, malaise, 10 and weight loss or, occasionally, more severe symptoms.

The progression from fibrosis, an early consequence of liver disease, to cirrhosis, and the specific histologic morphology that characterizes cirrhosis depend on the extent of injury, the presence of continuing 15 damage, and the response of the liver to damage. The liver may be injured acutely and severely (e.g. necrosis with hepatitis), moderately over months or years (e.g. biliary tract obstruction and chronic active hepatitis), or modestly but continuously (e.g. alcohol abuse).

During the repair process, new vessels connecting the hepatic artery and portal vein to the hepatic venules form within the fibrous sheath that surrounds the surviving nodules of liver cells. These vessels restore the intrahepatic circulatory pathway, but 25 provide relatively low-volume, high-pressure drainage that is less efficient than normal and results in increased portal vein pressure (portal hypertension). cirrhosis is not static and its features depend on the disease activity and stage.

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As cirrhosis is the end stage of many forms of liver disease, many genes have been identified that can contribute to the development of cirrhosis. These include, e.g., the genes responsible for Wilson disease (Online Mendelian Inheritance of Man ("OMIM") 277900), type IV 35 qlycogen storage disease (OMIM 232500), galactosemia (OMIM

230400), and a deficiency of alpha-1-antitrypsin (OMIM 107400). There is substantial evidence, however, for as yet uncharacterized loci which cause cirrhosis.

For example, Iber and Maddrey, Prog. Liver Dis. 5 2: 290-302 (1965), reviewed 13 previously reported families and 8 new to this study, each with 2 or more affected members. They pointed out that, with a single exception, the multiple cases were in the same generation. Within a given family, the age of onset, clinical course, and biopsy 10 findings were very similar, but there were wide differences between families.

Kalra et al., Hum. Hered. 32:170-175 (1982) studied the families of 220 cases of Indian childhood cirrhosis and 70 families of age-matched controls. The 15 hypotheses of autosomal recessive, partial sex-linkage, and doubly recessive inheritance were found untenable and the authors concluded that multifactorial inheritance was most plausible. Lefkowitch et al., New Eng. J. Med. 307:271-277 (1982) described 4 white American sibs who died between 20 ages 4.5 and 6 years of cirrhosis that closely resembled that of the childhood cirrhosis of Asiatic Indians.

Another example of uncharacterized loci which cause cirrhosis are those related to the risk of alcoholism.

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Cloninger, Science 236:410-416 (1987), defined two separate types of alcoholism. According to these definitions, type 1 alcohol abuse has its usual onset after the age of 25 years and is characterized by severe psychological dependence and guilt. Type 1 occurs in both 30 men and women and requires both genetic and environmental factors to become manifest. By contrast, type 2 alcohol abuse has its onset before the age of 25; persons with this type of alcoholism are characterized by their inability to abstain from alcohol and by frequent aggressive and 35 antisocial behavior. Type 2 alcoholism is rarely found in

PCT/US01/00669 WO 01/57277

women and is much more heritable.

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Despite considerable effort to identify genes related to the risk of alcoholism, relatively few genes have been identified. Some of this work has suggested a 5 relationship between the metabolism of dopamine and alcoholism. Blum et al., J.A.M.A. 263:2055-2060 (1990) and Bolos et al., J.A.M.A. 264:3156-3160 (1990) investigated the relationship of the dopamine D2 receptor (DRD2; OMIM 126450) to alcoholism, but the sample size was small and 10 their results were inconclusive. However, Tiihonen et al., Molec. Psychiat. 4, 286-289 (1999), found a markedly higher frequency in a population of type 1 alcoholics of the low activity allele of the enzyme catechol-O-methyltransferase (COMT, OMIM 116790), which has a crucial role in the 15 metabolism of dopamine, suggesting a role for dopamine metabolism in increased risk of alcoholism. For a brief review of recent progress toward the identification of genes related to risk for alcoholism see Buck, Genome 9:927-928 (1998).

As another example, multiple genes have been shown to predispose to hyperlipoproteinemia or hyperlipidemia. Much attention has been focused on these disorders because there is a strong association of hyperlipidemia, especially hypercholesterolemia, with 25 development of coronary artery disease. Coronary artery disease accounts for at least 25% of all deaths in the United States. Coronary artery disease results when the arteries supplying the heart muscle become occluded by plaques composed of lipids like cholesterol, blood clotting 30 components and blood cells.

The major plasma lipids circulate bound to proteins as macromolecular complexes called lipoproteins. Although closely interrelated, the major lipoprotein classes - chylomicron, very-low-density lipoprotein (VLDL), 35 low-density lipoprotein (LDL), and high-density lipoprotein

(HDL) - are usually classified in terms of physicochemical properties (e.g., density after centrifugation). Chylomicrons, the largest lipoproteins, carry exogenous triglyceride from the intestine via the thoracic duct to 5 the venous system and into peripheral sites. VLDL carries endogenous triglyceride primarily from the liver to the same peripheral sites for storage or use. Lipases quickly degrade the triglyceride in VLDL to produce intermediate density lipoproteins (IDL) and within 2 to 6 h, IDL is 10 degraded further to generate LDL, which has a plasma halflife of 2 to 3 days. While the overall fate of LDL is unclear, the liver is responsible for removing approximately 70% and active receptor sites have been found on the surfaces of hepatocytes.

Several monogenic conditions that lead to elevated levels of one or more serum lipoproteins have been defined and the responsible gene identified, including, e.g., hyperlipoproteinemia type I (OMIM 238600), familial hypercholesterolemia (OMIM 143890), and familial defective 20 apolipoprotein B (OMIM 107730). However, in many cases the etiology is unknown and there is strong evidence for additional uncharacterized loci.

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For example, Zuliani et al., Arterioscler. Thromb. Vasc. Biol. 19:802-809 (1999) identified a 25 Sardinian family with a recessive form of hypercholesterolemia with the clinical features of familial hypercholesterolemia (OMIM 603813), and found that previously identified genes were not responsible for this disorder. They proposed that in this new lipid disorder, a 30 recessive defect causes a selective impairment of the LDL receptor function in the liver. Ciccarese et al., Am. J. Hum. Genet. 66:453-460 (2000) recently mapped this novel disease locus.

Another example is designated familial combined 35 hyperlipidemia (OMIM 144250) which affects approximately 1-

PCT/US01/00669 WO 01/57277

2% of the population in the Western world. This disorder can have its basis in mutation in several novel genes, two of which have been mapped to chromosome 1 (Pajukanta et al., Nature Genet. 18:369-373 (1998)) and chromosome 11 5 (Aouizerat et al., Am. J. Hum. Genet. 65, 397-412 (1999)). The high frequency of this disorder suggests that most, if not all, hyperlipidemias are of multifactorial genetic etiology.

As yet a further example, primary schlerosing 10 cholangitis (PSC) is a disorder characterized by a patchy obliterative inflammatory fibrosis of the large bile ducts. Chronic inflammation leads to extensive bile duct strictures, cholestasis, and gradual progression to biliary cirrhosis. PSC occurs most often in young men and is 15 commonly associated with inflammatory bowel disease, especially ulcerative colitis. The onset is usually insidious, with gradual, progressive fatigue, pruritus, and jaundice. There is no specific therapy for sclerosing cholangitis, and liver transplantation is the only apparent 20 cure.

The etiology of PSC is not known, but both genetic and immunologic abnormalities have been implicated. However, the frequency of HLA-B8 and HLA-DT2, which are associated with a number of autoimmune diseases, is higher 25 in PSC than normal individuals. Prochazka et al., New Eng. J. Med. 322:1842-1844 (1990) found that 100% of 29 patients with primary sclerosing cholangitis carried the HLA-DRw52a antigen, which is normally present in 35% of the population.

As a still further example, sarcoidosis is a disease of unknown cause characterized by non-caseating granulomas in one or more organ systems. These granulomas may resolve completely or proceed to fibrosis. The disorder is systemic, but the liver is affected in approximately 75% of cases. Sarcoidosis occurs mainly in persons aged 20 to

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40 yr and is most common in Northern Europeans and American blacks. The lifetime risk of developing sarcoidosis is particularly high among Swedish men (1.15%), Swedish women (1.6%), and African Americans (2.4%).

The much greater frequency in African Americans 5 relative to the United States population overall suggests a genetic contribution to etiology. Early research studying familial aggregation indicated that the disease may have a nongenetic basis because the family pattern did not conform 10 to a simple Mendelian mode of inheritance (Allison, Sth. Med. J. 57: 27-32 (1964)). However, Headings et al., Ann. N.Y. Acad. Sci. 278:377-385 (1976) favored multifactorial genetic inheritance of susceptibility. Nowack et al., Arch. Intern. Med. 147:481-483 (1987), found an unusually 15 high frequency of HLA-DR5 in a study of 440 patients with sarcoidosis in Marburg, Germany. They also concluded that the role of an environmental or infectious agent triggering sarcoidosis cannot be envisaged without considering genetically linked cofactors.

Other significant diseases of liver are also 20 believed to have a genetic, typically polygenic, etiologic component. These diseases include, e.g., primary biliary cirrhosis, Zellweger syndrome, cholestasis-lymphedema syndrome, Alstrom syndrome, primary pulmonary 25 hypertension, Berardinelli-Seip congenital lipodystrophy, iron overload in Africa, neonatal cholestatic hepatitis, autosomal recessive KID syndrome, familial hypotransferrinemia, type I congenital dyserythropoietic anemia, porphyria variegata, Finnish lactic acidosis with 30 hepatic hemosiderosis, Rotor syndrome, essential hypertension, ARC syndrome, type II conjugated hyperbilirubinemia, Lambert syndrome, ichthyosis congenita with biliary atresia, Kabuki make-up syndrome, Meckel syndrome, cerebral aneurysm-cirrhosis syndrome, glycogen 35 storage diseases, polycystic kidney and hepatic disease,

isolated Caroli disease, trisomy 18-like syndrome, Osler-Rendu-Weber syndrome 3, fatal intrahepatic cholestasis, Coach syndrome, type C Niemann-Pick disease, hereditary persistence of fetal hemoglobin and hepatocellular cancer.

Altered responses to a variety of infectious agents that target the liver, especially acute viral hepatitis, have also been shown or are suspected to have genetic bases or contributions. In addition to differential susceptibility to primary infectious agents, these altered responses include predisposition to complicating conditions following contact with particular infectious agents. These include, e.g., development of hepatocellular carcinoma 2 correlated with Hepatitis B infection, and severe hepatic fibrosis following

Schistosoma mansoni infection.

The central role of the liver in drug metabolism results in exposure of this organ to a large variety of potentially toxic chemical agents and metabolites. These include naturally occurring plant alkaloids and mycotoxins, industrial chemicals, and, additionally, pharmacologic agents used in treating disease. The range of manifestations of toxin- and drug-induced liver disease are virtually as broad as the range of acute and chronic disorders and have also been shown or suspected to have genetic bases or contributions.

Such interactions between drugs and genotype have been shown in the response, e.g., to the anticonvulsant phenytoin, which can cause severe hepatitis-like disease in individuals who are impaired in the ability to detoxify a metabolite of phenytoin in the liver, and in the response to the drug sodium valproate, which can produce severe hepatotoxicity in certain individuals. The abnormal responses to both of these drugs are believed to be influenced by underlying genetic factors.

The human genome-derived single exon nucleic acid

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probes and microarrays of the present invention are useful for predicting, diagnosing, grading, staging, monitoring and prognosing diseases of human liver, particularly those diseases with polygenic etiology. With each of the single 5 exon probes described herein shown to be expressed at detectable levels in human liver, and with about 2/3 of the probes identifying novel genes, the single exon microarrays of the present invention provide exceptionally high informational content for such studies.

For example, diagnosis (including differential diagnosis among clinically indistinguishable disorders, such as cirrhosis), staging, and/or grading of a disease can be based upon the quantitative relatedness of a patient gene expression profile to one or more reference expression 15 profiles known to be characteristic of a given liver disease, or to specific grades or stages thereof.

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In one embodiment, the patient gene expression profile is generated by hybridizing nucleic acids obtained directly or indirectly from transcripts expressed in the 20 patient's liver to the genome-derived single exon microarray of the present invention. Reference profiles are obtained similarly, using nucleic acids obtained directly or indirectly from transcripts expressed by liver of individuals with known liver disease. Methods for 25 quantitatively relating gene expression profiles, without regard to the function of the protein encoded by the gene, are disclosed in WO 99/58720, incorporated herein by reference in its entirety.

In another approach, the genome-derived single 30 exon probes and microarrays of the present invention can be used to interrogate genomic DNA, rather than pools of expressed message; this latter approach permits predisposition to and/or prognosis of liver disease to be assessed through the massively parallel determination of 35 altered copy number, deletion, or mutation in the patient's

genome of exons known to be expressed in human liver. The algorithms set forth in WO 99/58720 can be applied to such genomic profiles without regard to the function of the protein encoded by the interrogated gene.

The utility is specific to the probe; at sufficiently high hybridization stringency, which stringencies are well known in the art — see Ausubel et al. and Maniatis et al. — each probe reports the level of expression of message specifically containing that ORF.

It should be appreciated, however, that the probes of the present invention, for which expression in the Fetal liver has been demonstrated are useful for both measurement in the Fetal liver and for survey of expression in other tissues.

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Significant among such advantages is the presence of probes for novel genes.

As mentioned above and further detailed in
Examples 1 and 2, the methods described enable ORFs which
are not present in existing expression databases to be

20 identified. And the fewer the number of tissues in which
the ORF can be shown to be expressed, the more likely the
ORF will prove to be part of a novel gene: as further
discussed in Example 2, ORFs whose expression was
measurable in only a single of the tested tissues were

25 represented in existing expression databases at a rate of
only 11%, whereas 36% of ORFs whose expression was
measurable in 9 tissues were present in existing expression
databases, and fully 45% of those ORFs expressed in all ten
tested tissues were present in existing expressed sequence
30 databases.

Either as tools for measuring gene expression or tools for surveying gene expression, the genome-derived single exon probes of the present invention have significant advantages over the cDNA or EST-based probes that are currently available for achieving these utilities.

The genome-derived single exon probes of the present invention are useful in constructing genome-derived single exon microarrays; the genome-derived single exon microarrays, in turn, are useful devices for measuring and for surveying gene expression in the human.

Gene expression analysis using microarrays — conventionally using microarrays having probes derived from expressed message — is well-established as useful in the biological research arts (see Lockhart et al. Nature 405, 827-836).

Microarrays have been used to determine gene expression profiles in cells in response to drug treatment (see, for example, Kaminski et al., "Global Analysis of Gene Expression in Pulmonary Fibrosis Reveals Distinct 15 Programs Regulating Lung Inflammation and Fibrosis, " Proc. Natl. Acad. Sci. USA 97(4):1778-83 (2000); Bartosiewicz et al., "Development of a Toxicological Gene Array and Quantitative Assessment of This Technology, " Arch. Biochem. Biophys. 376(1):66-73 (2000)), viral infection (see for 20 example, Geiss et al., "Large-scale Monitoring of Host Cell Gene Expression During HIV-1 Infection Using cDNA Microarrays, " Virology 266(1):8-16 (2000)) and during cell processes such as differentiation, senescence and apoptosis (see, for example, Shelton et al., "Microarray Analysis of 25 Replicative Senescence, " Curr. Biol. 9(17):939-45 (1999); Voehringer et al., "Gene Microarray Identification of Redox and Mitochondrial Elements That Control Resistance or Sensitivity to Apoptosis, " Proc. Natl. Acad. Sci. USA 97(6):2680-5 (2000)).

Microarrays have also been used to determine abnormal gene expression in diseased tissues (see, for example, Alon et al., "Broad Patterns of Gene Expression Revealed by Clustering Analysis of Tumor and Normal Colon Tissues Probed by Oligonucleotide Arrays," Proc. Natl.

35 Acad. Sci. USA 96(12):6745-50 (1999); Perou et al.,

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"Distinctive Gene Expression Patterns in Human Mammary
Epithelial Cells and Breast Cancers, Proc. Natl. Acad. Sci.
USA 96(16):9212-7 (1999); Wang et al., "Identification of
Genes Differentially Over-expressed in Lung Squamous Cell

Carcinoma Using Combination of cDNA Subtraction and
Microarray Analysis," Oncogene 19(12):1519-28 (2000);
Whitney et al., "Analysis of Gene Expression in Multiple
Sclerosis Lesions Using cDNA Microarrays," Ann. Neurol.
46(3):425-8 (1999)), in drug discovery screens (see, for
example, Scherf et al., "A Gene Expression Database for the
Molecular Pharmacology of Cancer," Nat. Genet. 24(3):236-44
(2000)) and in diagnosis to determine appropriate treatment
strategies (see, for example, Sgroi et al., "In vivo Gene
Expression Profile Analysis of Human Breast Cancer

Progression," Cancer Res. 59(22):5656-61 (1999)).

In microarray-based gene expression screens of pharmacological drug candidates upon cells, each probe provides specific useful data. In particular, it should be appreciated that even those probes that show no change in expression are as informative as those that do change, serving, in essence, as negative controls.

For example, where gene expression analysis is used to assess toxicity of chemical agents on cells, the failure of the agent to change a gene's expression level is evidence that the drug likely does not affect the pathway of which the gene's expressed protein is a part.

Analogously, where gene expression analysis is used to assess side effects of pharmacological agents — whether in lead compound discovery or in subsequent screening of lead compound derivatives — the inability of the agent to alter a gene's expression level is evidence that the drug does not affect the pathway of which the gene's expressed protein is a part.

WO 99/58720 provides methods for quantifying the relatedness of a first and second gene expression profile

and for ordering the relatedness of a plurality of gene expression profiles. The methods so described permit useful information to be extracted from a greater percentage of the individual gene expression measurements from a microarray than methods previously used in the art.

Other uses of microarrays are described in Gerhold et al., Trends Biochem. Sci. 24(5):168-173 (1999) and Zweiger, Trends Biotechnol. 17(11):429-436 (1999); Schena et al.

The invention particularly provides genomederived single-exon probes known to be expressed in Fetal liver. The individual single exon probes can be provided in the form of substantially isolated and purified nucleic acid, typically, but not necessarily, in a quantity sufficient to perform a hybridization reaction.

Such nucleic acid can be in any form directly hybridizable to the message that contains the probe's ORF, such as double stranded DNA, single-stranded DNA complementary to the message, single-stranded RNA complementary to the message, or chimeric DNA/RNA molecules so hybridizable. The nucleic acid can alternatively or additionally include either nonnative nucleotides, alternative internucleotide linkages, or both, so long as complementary binding can be obtained. For example, probes can include phosphorothioates, methylphosphonates, morpholino analogs, and peptide nucleic acids (PNA), as are described, for example, in U.S. Patent Nos. 5,142,047; 5,235,033; 5,166,315; 5,217,866; 5,184,444; 5,861,250.

Usefully, however, such probes are provided in a form and quantity suitable for amplification, where the amplified product is thereafter to be used in the hybridization reactions that probe gene expression.

Typically, such probes are provided in a form and quantity suitable for amplification by PCR or by other well known amplification technique. One such technique additional to

PCR is rolling circle amplification, as is described, inter alia, in U.S. Patent Nos. 5,854,033 and 5,714,320 and international patent publications WO 97/19193 and WO 00/15779. As is well understood, where the probes are to be provided in a form suitable for amplification, the range of nucleic acid analogues and/or internucleotide linkages will be constrained by the requirements and nature of the amplification enzyme.

Where the probe is to be provided in form

10 suitable for amplification, the quantity need not be sufficient for direct hybridization for gene expression analysis, and need be sufficient only to function as an amplification template, typically at least about 1, 10 or 100 pg or more.

packaged with amplification primers, either in a single composition that comprises probe template and primers, or in a kit that comprises such primers separately packaged therefrom. As earlier mentioned, the ORF-specific 5' primers used for genomic amplification can have a first common sequence added thereto, and the ORF-specific 3' primers used for genomic amplification can have a second, different, common sequence added thereto, thus permitting, in this embodiment, the use of a single set of 5' and 3' primers to amplify any one of the probes. The probe composition and/or kit can also include buffers, enzyme, etc., required to effect amplification.

As mentioned earlier, when intended for use on a genome-derived single exon microarray of the present invention, the genome-derived single exon probes of the present invention will typically average at least about 100, 200, 300, 400 or 500 bp in length, including (and typically, but not necessarily centered about) the ORF. Furthermore, when intended for use on a genome-derived single exon microarray of the present invention, the

genome-derived single exon probes of the present invention will typically not contain a detectable label.

When intended for use in solution phase hybridization, however - that is, for use in a 5 hybridization reaction in which the probe is not first bound to a support substrate (although the target may indeed be so bound) - length constraints that are imposed in microarray-based hybridization approaches will be relaxed, and such probes will typically be labeled.

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In such case, the only functional constraint that dictates the minimum size of such probe is that each such probe must be capable of specifically identifying in a hybridization reaction the exon from which it is drawn. In theory, a probe of as little as 17 nucleotides is capable 15 of uniquely identifying its cognate sequence in the human genome. For hybridization to expressed message - a subset of target sequence that is much reduced in complexity as compared to genomic sequence - even fewer nucleotides are required for specificity.

Therefore, the probes of the present invention 20 can include as few as 20, 25 or 50 bp or ORF, or more. In particular embodiments, the ORF sequences are given in SEQ ID NOS. 12,674 - 25,129, respectively, for probe SEQ ID NOS. 1 - 12,673. The minimum amount of ORF required to be 25 included in the probe of the present invention in order to provide specific signal in either solution phase or microarray-based hybridizations can readily be determined for each of ORF SEQ ID NOS. 12,674 - 25,129 individually by routine experimentation using standard high stringency 30 conditions.

Such high stringency conditions are described, inter alia, in Ausubel et al. and Maniatis et al. For microarray-based hybridization, standard high stringency conditions can usefully be 50% formamide, 5% SSC, 0.2 $\mu g/\mu l$ 35 poly(dA), 0.2 μ g/ μ l human cot1 DNA, and 0.5 % SDS, in a

humid oven at 42°C overnight, followed by successive washes of the microarray in 1% SSC, 0.2% SDS at 55°C for 5 minutes, and then 0.1% SSC, 0.2% SDS, at 55°C for 20 minutes. For solution phase hybridization, standard high 5 stringency conditions can usefully be aqueous hybridization at 65°C in 6X SSC. Lower stringency conditions, suitable for cross-hybridization to mRNA encoding structurally- and functionally-related proteins, can usefully be the same as the high stringency conditions but with reduction in 10 temperature for hybridization and washing to room temperature (approximately 25°C).

When intended for use in solution phase hybridization, the maximum size of the single exon probes of the present invention is dictated by the proximity of 15 other expressed exons in genomic DNA: although each single exon probe can include intergenic and/or intronic material contiguous to the ORF in the human genome, each probe of the present invention will include portions of only one expressed exon.

Thus, each single exon probe will include no more than about 25 kb of contiguous genomic sequence, more typically no more than about 20 kb of contiquous genomic sequence, more usually no more than about 15 kb, even more usually no more than about 10 kb. Usually, probes that are 25 maximally about 5 kb will be used, more typically no more than about 3 kb.

20

It will be appreciated that the Sequence Listing appended hereto presents, by convention, only that strand of the probe and ORF sequence that can be directly 30 translated reading from 5' to 3' end. As would be well understood by one of skill in the art, single stranded probes must be complementary in sequence to the ORF as present in an mRNA; it is well within the skill in the art to determine such complementary sequence. It will further 35 be understood that double stranded probes can be used in

both solution-phase hybridization and microarray-based hybridization if suitably denatured.

Thus, it is an aspect of the present invention to provide single-stranded nucleic acid probes that have

5 sequence complementary to those described herein above and below, and double-stranded probes one strand of which has sequence complementary to the probes described herein.

The probes can, but need not, contain intergenic and/or intronic material that flanks the ORF, on one or both sides, in the same linear relationship to the ORF that the intergenic and/or intronic material bears to the ORF in genomic DNA. The probes do not, however, contain nucleic acid derived from more than one expressed ORF.

And when intended for use in solution

hybridization, the probes of the present invention can
usefully have detectable labels. Nucleic acid labels are
well known in the art, and include, inter alia, radioactive
labels, such as ³H, ³²P, ³³P, ³⁵S, ¹²⁵I, ¹³¹I; fluorescent
labels, such as Cy3, Cy5, Cy5.5, Cy7, SYBR®

Green and other labels described in Haugland,

Handbook of Fluorescent Probes and Research Chemicals, 7th
ed., Molecular Probes Inc., Eugene, OR (2000), or
fluorescence resonance energy transfer tandem conjugates
thereof; labels suitable for chemiluminescent and/or
enhanced chemiluminescent detection; labels suitable for
ESR and NMR detection; and labels that include one member
of a specific binding pair, such as biotin, digoxigenin, or
the like.

The probes, either in quantity sufficient for 30 hybridization or sufficient for amplification, can be provided in individual vials or containers.

Alternatively, such probes can usefully be packaged as a plurality of such individual genome-derived single exon probes.

When provided as a collection of plural

individual probes, the probes are typically made available in amplifiable form in a spatially-addressable ordered set, typically one per well of a microtiter dish. Although a 96 well microtiter plate can be used, greater efficiency is obtained using higher density arrays.

If, as earlier mentioned, the ORF-specific
5' primers used for genomic amplification had a first
common sequence added thereto, and the ORF-specific 3'
primers used for genomic amplification had a second,
different, common sequence added thereto, a single set of
5' and 3' primers can be used to amplify all of the probes
from the amplifiable ordered set.

Such collections of genome-derived single exon probes can usefully include a plurality of probes chosen for the common attribute of expression in the human Fetal liver.

In such defined subsets, typically at least 50, 60, 75, 80, 85, 90 or 95% or more of the probes will be chosen by their expression in the defined tissue or cell type.

The single exon probes of the present invention, as well as fragments of the single exon probes comprising selectively hybridizable portions of the probe ORF, can be used to obtain the full length cDNA that includes the ORF by (i) screening of cDNA libraries; (ii) rapid amplification of cDNA ends ("RACE"); or (iii) other conventional means, as are described, inter alia, in Ausubel et al. and Maniatis et al.

It is another aspect of the present invention to provide genome-derived single exon nucleic acid microarrays useful for gene expression analysis, where the term "microarray" has the meaning given in the definitional section of this description, supra.

The invention particularly provides genome-35 derived single-exon nucleic acid microarrays comprising a

plurality of probes known to be expressed in human Fetal liver. In preferred embodiments, the present invention provides human genome-derived single exon microarrays comprising a plurality of probes drawn from the group consisting of SEQ ID NOS.: 1 - 12,673.

When used for gene expression analysis, the genome-derived single exon microarrays provide greater physical informational density than do the genome-derived single exon microarrays that have lower percentages of 10 probes known to be expressed commonly in the tested tissue. At a fixed probe density, for example, a given microarray surface area of the defined subset genome-derived single exon microarray can yield a greater number of expression measurements. Alternatively, at a given probe density, the 15 same number of expression measurements can be obtained from a smaller substrate surface area. Alternatively, at a fixed probe density and fixed surface area, probes can be provided redundantly, providing greater reliability in signal measurement for any given probe. Furthermore, with 20 a higher percentage of probes known to be expressed in the assayed tissue, the dynamic range of the detection means can be adjusted to reveal finer levels discrimination among the levels of expression.

Although particularly described with respect to

their utility as probes of gene expression, particularly as probes to be included on a genome-derived single exon microarray, each of the nucleic acids having SEQ ID NOS.: 1 - 12,673 contains an open-reading frame, set forth respectively in SEQ ID NOS.: 12,674 - 25,129, that encodes a protein domain. Thus, each of SEQ ID NOS. 1 - 12,673 can be used, or that portion thereof in SEQ ID NOS. 12,674 - 25,129 used, to express a protein domain by standard in vitro recombinant techniques. See Ausubel et al. and Maniatis et al.

Additionally, kits are available commercially

that readily permit such nucleic acids to be expressed as protein in bacterial cells, insect cells, or mammalian cells, as desired (e.g., HAT™ Protein Expression & Purification System, ClonTech Laboratories, Palo Alto, CA; Adeno-X™ Expression System, ClonTech Laboratories, Palo Alto, CA; Protein Fusion & Purification (pMAL™) System, New England Biolabs, Beverley, MA)

Furthermore, shorter peptides can be chemically synthesized using commercial peptide synthesizing equipment and well known techniques. Procedures are described, inter alia, in Chan et al. (eds.), Fmoc Solid Phase Peptide Synthesis: A Practical Approach (Practical Approach Series, (Paper)), Oxford Univ. Press (March 2000) (ISBN: 0199637245); Jones, Amino Acid and Peptide Synthesis (Oxford Chemistry Primers, No 7), Oxford Univ. Press (August 1992) (ISBN: 0198556683); and Bodanszky, Principles of Peptide Synthesis (Springer Laboratory), Springer Verlag (December 1993) (ISBN: 0387564314).

It is, therefore, another aspect of the invention
to provide peptides comprising an amino acid sequence
translated from SEQ ID NOS: 12,674 - 25,129. Such amino
acid sequences are set out in SEQ ID NOS: 25,130 - 37,156.
Any such recombinantly-expressed or synthesized peptide of
at least 8, and preferably at least about 15, amino acids,
can be conjugated to a carrier protein and used to generate
antibody that recognizes the peptide. Thus, it is a
further aspect of the invention to provide peptides that
have at least 8, preferably at least 15, consecutive amino
acids.

30

The following examples are offered by way of illustration and not by way of limitation.

EXAMPLE_1

35 Preparation of Single Exon Microarrays from ORFs Predicted

in Human Genomic Sequence

Bioinformatics Results

All human BAC sequences in fewer than 10 pieces

5 that had been accessioned in a five month period
immediately preceding this study were downloaded from
GenBank. This corresponds to ~2200 clones, totaling ~350
MB of sequence, or approximately 10% of the human genome.

After masking repetitive elements using the

10 program CROSS_MATCH, the sequence was analyzed for open
reading frames using three separate gene finding programs.
The three programs predict genes using independent
algorithmic methods developed on independent training sets:
GRAIL uses a neural network, GENEFINDER uses a hidden

15 Markoff model, and DICTION, a program proprietary to
Genetics Institute, operates according to a different
heuristic. The results of all three programs were used to
create a prediction matrix across the segment of genomic
DNA.

The three gene finding programs yielded a range of results. GRAIL identified the greatest percentage of genomic sequence as putative coding region, 2% of the data analyzed. GENEFINDER was second, calling 1%, and DICTION yielded the least putative coding region, with 0.8% of genomic sequence called as coding region.

The consensus data were as follows. GRAIL and GENEFINDER agreed on 0.7% of genomic sequence, GRAIL and DICTION agreed on 0.5% of genomic sequence, and the three programs together agreed on 0.25% of the data analyzed.

That is, 0.25% of the genomic sequence was identified by all three of the programs as containing putative coding region.

ORFs predicted by any two of the three programs
("consensus ORFs") were assorted into "gene bins" using two
criteria: (1) any 7 consecutive exons within a 25 kb window

were placed together in a bin as likely contributing to a single gene, and (2) all ORFs within a 25 kb window were placed together in a bin as likely contributing to a single gene if fewer than 7 exons were found within the 25 kb window.

PCR

The largest ORF from each gene bin that did not span repetitive sequence was then chosen for amplification, as were all consensus ORFs longer than 500 bp. This method approximated one exon per gene; however, a number of genes were found to be represented by multiple elements.

Previously, we had determined that DNA fragments fewer than 250 bp in length do not bind well to the aminomodified glass surface of the slides used as support substrate for construction of microarrays; therefore, amplicons were designed in the present experiments to approximate 500 bp in length.

Accordingly, after selecting the largest ORF per gene bin, a 500 bp fragment of sequence centered on the ORF was passed to the primer picking software, PRIMER3 (available online for use at http://www-genome.wi.mit.edu/cgi-bin/primer/). A first additional sequence was commonly added to each ORF-unique 5' primer, and a second, different, additional sequence was commonly added to each ORF-unique 3' primer, to permit subsequent reamplification of the amplicon using a single set of "universal" 5' and 3' primers, thus immortalizing the amplicon. The addition of universal priming sequences also facilitates sequence verification, and can be used to add a cloning site should some ORFs be found to warrant further study.

The ORFs were then PCR amplified from genomic DNA, verified on agarose gels, and sequenced using the universal primers to validate the identity of the amplicon

to be spotted in the microarray.

Primers were supplied by Operon Technologies

(Alameda, CA). PCR amplification was performed by standard techniques using human genomic DNA (Clontech, Palo Alto,

5 CA) as template. Each PCR product was verified by SYBR® green (Molecular Probes, Inc., Eugene, OR) staining of agarose gels, with subsequent imaging by Fluorimager (Molecular Dynamics, Inc., Sunnyvale, CA). PCR amplification was classified as successful if a single band appeared.

The success rate for amplifying ORFs of interest directly from genomic DNA using PCR was approximately 75%. FIG. 5 graphs the distribution of predicted ORF (exon) length and distribution of amplified PCR products, with ORF length shown in red and PCR product length shown in blue (which may appear black in the figure). Although the range of ORF sizes is readily seen to extend to beyond 900 bp, the mean predicted exon size was only 229 bp, with a median size of 150 bp (n=9498). With an average amplicon size of 475 ± 25 bp, approximately 50% of the average PCR amplification product contained predicted coding region, with the remaining 50% of the amplicon containing either intron, intergenic sequence, or both.

Using a strategy predicated on amplifying about 500 bp, it was found that long exons had a higher PCR failure rate. To address this, the bioinformatics process was adjusted to amplify 1000, 1500 or 2000 bp fragments from exons larger than 500 bp. This improved the rate of successful amplification of exons exceeding 500 bp, constituting about 9.2% of the exons predicted by the gene finding algorithms.

Approximately 75% of the probes disposed on the array (90% of those that successfully PCR amplified) were sequence-verified by sequencing in both the forward and reverse direction using MegaBACE sequencer (Molecular

Dynamics, Inc., Sunnyvale, CA), universal primers, and standard protocols.

Some genomic clones (BACs) yielded very poor PCR and sequencing results. The reasons for this are unclear, but may be related to the quality of early draft sequence or the inclusion of vector and host contamination in some submitted sequence data.

Although the intronic and intergenic material flanking coding regions could theoretically interfere with hybridization during microarray experiments, subsequent empirical results demonstrated that differential expression ratios were not significantly affected by the presence of noncoding sequence. The variation in exon size was similarly found not to affect differential expression ratios significantly; however, variation in exon size was observed to affect the absolute signal intensity (data not shown).

The 350 MB of genomic DNA was, by the above-described process, reduced to 9750 discrete probes, which were spotted in duplicate onto glass slides using commercially available instrumentation (MicroArray GenII Spotter and/or MicroArray GenIII Spotter, Molecular Dynamics, Inc., Sunnyvale, CA). Each slide additionally included either 16 or 32 E. coli genes, the average hybridization signal of which was used as a measure of background biological noise.

Each of the probe sequences was BLASTed against the human EST data set, the NR data set, and SwissProt GenBank (May 7, 1999 release 2.0.9).

30

One third of the probe sequences (as amplified) produced an exact match (BLAST Expect ("E") values less than 1 e⁻¹⁰⁰) to either an EST (20% of sequences) or a known mRNA (13% of sequences). A further 22% of the probe sequences showed some homology to a known EST or mRNA (BLAST E values from 1 e⁻⁵ to 1 e⁻⁹⁹). The remaining 45% of

the probe sequences showed no significant sequence homology to any expressed, or potentially expressed, sequences present in public databases.

All of the probe sequences (as amplified) were
then analyzed for protein similarities with the SwissProt
database using BLASTX, Gish et al., Nature Genet. 3:266
(1993). The predicted functional breakdowns of the 2/3 of
probes identical or homologous to known sequences are
presented in Table 1.

10

Table 1

	-		
Function	of Predic	ted ORFs As	Deduced From Comparative
Sequence	Analysis		
Total	V6 chip	V7 chip	Function Predicted from
			Comparative Sequence
			Analysis
211	96	115	Receptor
120	43	77	Zinc Finger
30	11	19	Homeobox
25	9	16	Transcription Factor
17	11	7	Transcription .
118	57	61	Structural
95	39	56	Kinase
36	18	18	Phosphatase
83	31	52	Ribosomal
45	19	26	Transport
21	17	14	Growth Factor
17	12	5	Cytochrome
50	33	17	Channel
1	1		

As can be seen, the two most common types of genes were transcription factors and receptors, making up 2.2% and 1.8% of the arrayed elements, respectively.

EXAMPLE 2

Gene Expression Measurements From Genome-Derived Single 5 Exon Microarrays

The two genome-derived single exon microarrays prepared according to Example 1 were hybridized in a series 10 of simultaneous two-color fluorescence experiments to (1) Cy3-labeled cDNA synthesized from message drawn individually from each of brain, heart, liver, fetal liver, placenta, lung, bone marrow, HeLa, BT 474, or HBL 100 cells, and (2) Cy5-labeled cDNA prepared from message 15 pooled from all ten tissues and cell types, as a control in each of the measurements. Hybridization and scanning were carried out using standard protocols and Molecular Dynamics equipment.

Briefly, mRNA samples were bought from commercial 20 sources (Clontech, Palo Alto, CA and Amersham Pharmacia Biotech (APB)). Cy3-dCTP and Cy5-dCTP (both from APB) were incorporated during separate reverse transcriptions of 1 μg of polyA* mRNA performed using 1 µg oligo(dT)12-18 primer and 2 μg random 9mer primers as follows. After heating to 25 70°C, the RNA:primer mixture was snap cooled on ice. After snap cooling on ice, added to the RNA to the stated final concentration was: 1X Superscript II buffer, 0.01 M DTT, 100 μ M dATP, 100 μ M dGTP, 100 μ M dTTP, 50 μ M dCTP, 50 μ M Cy3-dCTP or Cy5-dCTP 50 μM , and 200 U Superscript II 30 enzyme. The reaction was incubated for 2 hours at 42°C. After 2 hours, the first strand cDNA was isolated by adding 1 U Ribonuclease H, and incubating for 30 minutes at 37°C. The reaction was then purified using a Qiagen PCR cleanup column, increasing the number of ethanol washes to 5.

35 Probe was eluted using 10 mM Tris pH 8.5.

Using a spectrophotometer, probes were measured for dye incorporation. Volumes of both Cy3 and Cy5 cDNA corresponding to 50 pmoles of each dye were then dried in a Speedvac, resuspended in 30 µl hybridization solution containing 50% formamide, 5X SSC, 0.2 µg/µl poly(dA), 0.2 µg/µl human cot1 DNA, and 0.5 % SDS.

Hybridizations were carried out under a coverslip, with the array placed in a humid oven at 42°C overnight. Before scanning, slides were washed in 1X SSC, 0.2% SDS at 55°C for 5 minutes, followed by 0.1X SSC, 0.2% SDS, at 55°C for 20 minutes. Slides were briefly dipped in water and dried thoroughly under a gentle stream of nitrogen.

Slides were scanned using a Molecular Dynamics

Gen3 scanner, as described. Schena (ed.), Microarray

Biochip: Tools and Technology, Eaton Publishing

Company/BioTechniques Books Division (2000) (ISBN:

1881299376).

Although the use of pooled cDNA as a reference

20 permitted the survey of a large number of tissues, it
attenuates the measurement of relative gene expression,
since every highly expressed gene in the tissue/cell typespecific fluorescence channel will be present to a level of
at least 10% in the control channel. Because of this fact,

25 both signal and expression ratios (the latter hereinafter,
"expression" or "relative expression") for each probe were
normalized using the average ratio or average signal,
respectively, as measured across the whole slide.

Data were accepted for further analysis only when signal was at least three times greater than biological noise, the latter defined by the average signal produced by the E. coli control genes.

The relative expression signal for these probes was then plotted as function of tissue or cell type, and is presented in FIG. 6.

FIG. 6 shows the distribution of expression across a panel of ten tissues. The graph shows the number of sequence-verified products that were either not expressed ("0"), expressed in one or more but not all tested tissues ("1" - "9"), and expressed in all tissues tested ("10").

Of 9999 arrayed elements on the two microarrays (including positive and negative controls and "failed" products), 2353 (51%) were expressed in at least one tissue or cell type. Of the gene elements showing significant signal — where expression was scored as "significant" if the normalized Cy3 signal was greater than 1, representing signal 5-fold over biological noise (0.2) — 39% (991) were expressed in all 10 tissues. The next most common class (15%) consisted of gene elements expressed in only a single tissue.

The genes expressed in a single tissue were further analyzed, and the results of the analyses are compiled in FIG. 7.

FIG. 7A is a matrix presenting the expression of 20 all verified sequences that showed expression greater than 3 in at least one tissue. Each clone is represented by a column in the matrix. Each of the 10 tissues assayed is represented by a separate row in the matrix, and relative 25 expression of a clone in that tissue is indicated at the respective node by intensity of green shading, with the intensity legend shown in panel B. The top row of the matrix ("EST Hit") contains "bioinformatic" rather than "physical" expression data - that is, presents the results 30 returned by query of EST, NR and SwissProt databases using the probe sequence. The legend for "bioinformatic expression" (i.e., degree of homology returned) is presented in panel C. Briefly, white is known, black is novel, with gray depicting nonidentical with significant 35 homology (white: E values < 1e-100; gray: E values from 1e-

05 to 1e-99; black: E values > 1e-05).

As FIG. 7 readily shows, heart and brain were demonstrated to have the greatest numbers of genes that were shown to be uniquely expressed in the respective tissue. In brain, 200 uniquely expressed genes were identified; in heart, 150. The remaining tissues gave the following figures for uniquely expressed genes: liver, 100; lung, 70; fetal liver, 150; bone marrow, 75; placenta, 100; HeLa, 50; HBL, 100; and BT474, 50.

It was further observed that there were many more "novel" genes among those that were up-regulated in only one tissue, as compared with those that were down-regulated in only one tissue. In fact, it was found that ORFs whose expression was measurable in only a single of the tested tissues were represented in sequencing databases at a rate of only 11%, whereas 36% of the ORFs whose expression was measurable in 9 of the tissues were present in public databases. As for those ORFs expressed in all ten tissues, fully 45% were present in existing expressed sequence databases. These results are not unexpected, since genes expressed in a greater number of tissues have a higher likelihood of being, and thus of having been, discovered by EST approaches.

25 Comparison of Signal from Known and Unknown Genes

The normalized signal of the genes found to have high homology to genes present in the GenBank human EST database were compared to the normalized signal of those genes not found in the GenBank human EST database. The data are shown in FIG. 8.

FIG. 8 shows the normalized Cy3 signal intensity for all sequence-verified products with a BLAST Expect ("E") value of greater than 1e-30 (designated "unknown") upon query of existing EST, NR and SwissProt databases, and shows in blue the normalized Cy3 signal intensity for all

sequence-verified products with a BLAST Expect value of less than 1e-30 ("known"). Note that biological background noise has an averaged normalized Cy3 signal intensity of 0.2.

As expected, the most highly expressed of the 5 ORFs were "known" genes. This is not surprising, since very high signal intensity correlates with very commonlyexpressed genes, which have a higher likelihood of being found by EST sequence.

However, a significant point is that a large number of even the high expressers were "unknown". Since the genomic approach used to identify genes and to confirm their expression does not bias exons toward either the 3' or 5' end of a gene, many of these high expression genes 15 will not have been detected in an end-sequenced cDNA library.

The significant point is that presence of the gene in an EST database is not a prerequisite for incorporation into a genome-derived microarray, and 20 further, that arraying such "unknown" exons can help to assign function to as-yet undiscovered genes.

Verification of Gene Expression

10

To ascertain the validity of the approach 25 described above to identify genes from raw genomic sequence, expression of two of the probes was assayed using reverse transcriptase polymerase chain reaction (RT PCR) and northern blot analysis.

Two microarray probes were selected on the basis 30 of exon size, prior sequencing success, and tissue-specific gene expression patterns as measured by the microarray experiments. The primers originally used to amplify the two respective ORFs from genomic DNA were used in RT PCR against a panel of tissue-specific cDNAs (Rapid-Scan gene 35 expression panel 24 human cDNAs) (OriGene Technologies,

Inc., Rockville, MD).

Sequence AL079300 1 was shown by microarray hybridization to be present in cardiac tissue, and sequence AL031734 1 was shown by microarray experiment to be present 5 in placental tissue (data not shown). RT-PCR on these two sequences confirmed the tissue-specific gene expression as measured by microarrays, as ascertained by the presence of a correctly sized PCR product from the respective tissue type cDNAs.

Clearly, all microarray results cannot, and indeed should not, be confirmed by independent assay methods, or the high throughput, highly parallel advantages of microarray hybridization assays will be lost. However, in addition to the two RT-PCR results presented above, the 15 observation that 1/3 of the arrayed genes exist in expression databases provides powerful confirmation of the power of our methodology - which combines bioinformatic prediction with expression confirmation using genomederived single exon microarrays - to identify novel genes 20 from raw genomic data.

To verify that the approach further provides correct characterization of the expression patterns of the identified genes, a detailed analysis was performed of the microarrayed sequences that showed high signal in brain.

For this latter analysis, sequences that showed high (normalized) signal in brain, but which showed very low (normalized) signal (less than 0.5, determined to be biological noise) in all other tissues, were further studied. There were 82 sequences that fit these criteria, 30 approximately 2% of the arrayed elements. The 10 sequences showing the highest signal in brain in microarray hybridizations are detailed in Table 2, along with assigned function, if known or reasonably predicted.

35 Table 2

10

F	unction	of the Mos	st Highly	
Expressed G	enes Exp	ressed Onl	ly in Brain	า
Microarray	Normal	Homology	Gene Function	
Sequence	ized	on Ratio	to EST	as described by
Name	Signal		present	GenBank
	•		in	
			GenBank	
AP000217-1	5.2	+7.7	High	S-100 protein,
				b-chain, Ca ²⁺
				binding protein
				expressed in
				central nervous
				system
AP000047-1	2.3		High	Unknown
				Function
AC006548-9	1.7		High	Similar to '
,				mouse membrane
				glyco-protein
				M6, expressed
·				in central
				nervous system
AC007245-5	1.5		High	Similar to
				amphiphysin, a
				synaptic
				vesicle-
				associated
				protein. Ref 21
L44140-4	1.2	+2.0	High	Endothelial
			,	actin-binding
				protein found
				in nonmuscle
				filamin
, '	•	'	1	'

AC004689-9	1 3	1,2 6	Itiah	Protein
AC004689-9	1.2	+3.5	High	
				Phosphatase
				PP2A, neuronal/
				downregulates
				activated
				protein kinases
AL031657-1	1.2	+3.0	High	Unknown
				function/
				Contains the
				anhyrin motif,
				a common
				protein
;				sequence motif
AC009266-2	1.1	+3.7	Low	Low homology to
				the
				Synaptotagmin I
				protein in
				rat/present at
				low levels
		,		throughout rat
				brain
AP000086-1	1.0	+2.7	Low	Unknown, very
				poor homology
				to collagen
AC004689-3	1.0		High	Protein
				Phosphatase
			!	PP2A, neuronal/
				downregulates
		٠.		activated
		*		protein kinases

Of the ten sequences studied by these latter confirmatory approaches, eight were previously known. Of these eight, six had previously been reported to be

5 important in the central nervous system or brain. The exon

giving the highest signal (AP00217-1) was found to be the gene encoding an S100B Ca²⁺ binding protein, reported in the literature to be highly and uniquely expressed in the central nervous system. Heizmann, Neurochem. Res. 9:1097 (1997).

A number of the brain-specific probe sequences (including AC006548-9, AC009266-2) did not have homology to any known human cDNAs in GenBank but did show homology to rat and mouse cDNAs. Sequences AC004689-9 and AC004689-3 were both found to be phosphatases present in neurons (Millward et al., Trends Biochem. Sci. 24(5):186-191 (1999)). Two microarray sequences, AP000047-1 and AP000086-1 have unknown function, with AP000086-1 being absent from GenBank. Functionality can now be narrowed down to a role in the central nervous system for both of these genes, showing the power of designing microarrays in this fashion.

Next, the function of the chip sequences with the highest (normalized) signal intensity in brain, regardless of expression in other tissues, was assessed. In this latter analysis, we found expression of many more common genes, since the sequences were not limited to those expressed only in brain. For example, looking at the 20 highest signal intensity spots in brain, 4 were similar to tubulin (AC00807905; AF146191-2; AC007664-4; AF14191-2), 2 were similar to actin (AL035701-2; AL034402-1), and 6 were found to be homologous to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (AL035604-1; Z86090-1; AC006064-L, AC006064-K; AC035604-3; AC006064-L). These genes are often used as controls or housekeeping genes in microarray experiments of all types.

Other interesting genes highly expressed in brain were a ferritin heavy chain protein, which is reported in the literature to be found in brain and liver (Joshi et al., J. Neurol. Sci. 134(Suppl):52-56 (1995)), a result

duplicated with the array. Other highly expressed chip sequences included a translation elongation factor 10 (AC007564-4), a DEAD-box homolog (AL023804-4), and a Ychromosome RNA-binding motif (Chai et al., Genomics 5 49(2):283-89 (1998)) (AC007320-3). A low homology analog (AP00123-1/2) to a gene, DSCR1; thought to be involved in trisomy 21 (Down's syndrome), showed high expression in both brain and heart, in agreement with the literature (Fuentes et al., Mol. Genet. 4(10):1935-44 (1995)).

As a further validation of the approach, we selected the BAC AC006064 to be included on the array. This BAC was known to contain the GAPDH gene, and thus could be used as a control for the ORF selection process. The gene finding and exon selection algorithms resulted in 15 choosing 25 exons from BAC AC006064 for spotting onto the array, of which four were drawn from the GAPDH gene. Table 3 shows the comparison of the average expression ratio for the 4 exons from BAC006064 compared with the average expression ratio for 5 different dilutions of a 20 commercially available GAPDH cDNA (Clontech).

Table 3

Comparis	son of Expression R	atio, for each
tissue, of GAPDH		
	AC006064 (n = 4)	Control (n = 5)
Bone Marrow	-1.81 ± 0.11	-1.85 ± 0.08
Brain	-1.41 ± 0.11	-1.17 ± 0.05
BT474 .	1.85 ± 0.09	1.66 ± 0.12
Fetal Liver	-1.62 ± 0.07	-1.41 ± 0.05
HBL100	1.32 ± 0.05	2.64 ± 0.12
Heart	1.16 ± 0.09	1.56 ± 0.10
HeLa	1.11 ±0.06	1.30 ± 0.15
Liver	-1.62 ± 0.22	-2.07 ±

Lung	-4.95 ± 0.93	-3.75 ± 0.21
Placenta	-3.56 ± 0.25	-3.52 ± 0.43

Each tissue shows excellent agreement between the experimentally chosen exons and the control, again

5 demonstrating the validity of the present exon mining approach. In addition, the data also show the variability of expression of GAPDH within tissues, calling into question its classification as a housekeeping gene and utility as a housekeeping control in microarray

10 experiments.

EXAMPLE 3

Representation of Sequence and Expression Data as a "Mondrian"

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For each genomic clone processed for microarray as above-described, a plethora of information was accumulated, including full clone sequence, probe sequence within the clone, results of each of the three gene finding programs, EST information associated with the probe sequences, and microarray signal and expression for multiple tissues, challenging our ability to display the information.

Accordingly, we devised a new tool for visual display of the sequence with its attendant annotation which, in deference to its visual similarity to the paintings of Piet Mondrian, is hereinafter termed a "Mondrian". FIGS. 3 and 4 present the key to the information presented on a Mondrian.

FIG. 9 presents a Mondrian of BAC AC008172 (bases 25,000 to 130,000 shown), containing the carbamyl phosphate synthetase gene (AF154830.1). Purple background within the region shown as field 81 in FIG. 3 indicates all 37 known

exons for this gene.

As can be seen, GRAIL II successfully identified 27 of the known exons (73%), GENEFINDER successfully identified 37 of the known exons (100%), while DICTION identified 7 of the known exons (19%).

Seven of the predicted exons were selected for physical assay, of which 5 successfully amplified by PCR and were sequenced. These five exons were all found to be from the same gene, the carbamyl phosphate synthetase gene (AF154830.1).

The five exons were arrayed, and gene expression measured across 10 tissues. As is readily seen in the Mondrian, the five chip sequences on the array show identical expression patterns, elegantly demonstrating the reproducibility of the system.

FIG. 10 is a Mondrian of BAC AL049839. We selected 12 exons from this BAC, of which 10 successfully sequenced, which were found to form between 5 and 6 genes. Interestingly, 4 of the genes on this BAC are protease inhibitors. Again, these data elegantly show that exons 20 selected from the same gene show the same expression patterns, depicted below the red line. From this figure, it is clear that our ability to find known genes is very good. A novel gene is also found from 86.6 kb to 88.6 kb, 25 upon which all the exon finding programs agree. We are confident we have two exons from a single gene since they show the same expression patterns and the exons are proximal to each other. Backgrounds in the following colors indicate a known gene (top to bottom): 30 red = kallistatin protease inhibitor (P29622); purple = plasma serine protease inhibitor (P05154); turquoise = α 1 anti-chymotrypsin (P01011); mauve = 40S \ ribosomal protein (P08865). Note that chip sequence 8 and 12 did not sequence verify.

EXAMPLE 4

Genome-Derived Single Exon Probes Useful For Measuring Human Gene Expression

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The protocols set forth in Examples 1 and 2, supra, were applied to additional human genomic sequence as it became newly available in GenBank to identify unique exons in the human genome that could be shown to be 10 expressed at significant levels in Fetal liver tissue.

These unique exons are within longer probe sequences. Each probe was completely sequenced on both strands prior to its use on a genome-derived single exon microarray; sequencing confirms the exact chemical 15 structure of each probe. An added benefit of sequencing is that it placed us in possession of a set of single baseincremented fragments of the sequenced nucleic acid, starting from the sequencing primer 3' OH. (Since the single exon probes were first obtained by PCR amplification 20 from genomic DNA, we were of course additionally in possession of an even larger set of single base incremented fragments of each of the 12,673 single exon probes, each fragment corresponding to an extension product from one of the two amplification primers.)

The structures of the 12,673 unique single exon probes are clearly presented in the Sequence Listing as SEQ ID Nos.: 1 - 12,673. The 16 nt 5' primer sequence and 16 nt 3' primer sequence present on the amplicon are not included in the sequence listing. The sequences of the 30 exons present within each of these probes is presented in the Sequence Listing as SEQ ID Nos.: 12,674 - 25,129, respectively. It will be noted that some amplicons have more than one exon, some exons are contained in more than one amplicon.

35 As detailed in Example 2, expression was

demonstrated by disposing the amplicons as single exon probes on nucleic acid microarrays and then performing two-color fluorescent hybridization analysis; significant expression is based on a statistical confidence that the signal is significantly greater than negative biological control spots. The negative biological control is formed from spotted DNA sequences from a different species. Here, 32 sequences from E.Coli were spotted in duplicate to give a total of 64 spots.

10 For each hybridisation (each slide, each colour) the median value of the signal from all of the spots is determined. The normalised signal value is the arithmetic mean of the signal from duplicate spots divided by the population median.

15 Control spots are eliminated if there is more that a five-fold difference between each one of the duplicate spots raw signals.

The median of the signal from the remaining control spots is calculated and all subsequent calculations are done with normalised signals.

Control spots having a signal of greater than median + 2.4 (the value 2.4 is roughly 12 times the observed standard deviation of control spot populations) are eliminated. Spots with such high signals are considered to be "outliers".

The mean and standard deviation of the modified control spot populations are calculated.

The mean + 3x the standard deviation (mean + (3*SD)) is used as the signal threshold qualifier for that particular hybridisation. Thus, individual thresholds are determined for each channel and each hybridisation.

This means that, assuming that the data is distributed normally, there is a 99% confidence that any signal exceeding the threshold is significant.

35 The probes and their expression data are

presented in Table 4, set forth respectively in Example 5.

Example 5 presents the subset of probes that is significantly expressed in the human Fetal liver and thus presents the subset of probes that was recognized to be useful for measuring expression of their cognate genes in human Fetal liver tissue.

The sequence of each of the exon probes identified by SEQ ID NOS.: 12,674 to 25,129 was individually used as a BLAST (or, for SWISSPROT, BLASTX)

10 query to identify the most similar sequence in each of dbEST, SwissProt (BLASTX), and NR divisions of GenBank. Because the query sequences are themselves derived from genomic sequence in GenBank, only nongenomic hits from NR were scored.

15 The smallest in value of the BLAST (or BLASTX)
expect ("E") scores for each query sequence across the
three database divisions was used as a measure of the
"expression novelty" of the probe's ORF. Table 4 is sorted
in descending order based on this measure, reported as
20 "Most Similar (top) Hit BLAST E Value". Those sequences for
which no "Hit E Value" is listed are those exons which were
found to have no similar sequences.

As sorted, Table 4 thus lists its respective probes (by "AMPLICON SEQ ID NO.:" and additionally by the SEQ ID NO:. of the exon contained within the probe: "EXON SEQ ID NO.:") from least similar to sequences known to be expressed (i.e., highest BLAST E value), at the beginning of the table, to most similar to sequences known to be expressed (i.e., lowest BLAST E value), at the bottom of the table.

Table 4 further provides, for each listed probe, the accession number of the database sequence that yielded the "Most Similar (top) Hit BLAST E Value", along with the name of the database in which the database sequence is found ("Top Hit Database Source").

Table 4 further provides SEQ ID NOS. corresponding to the predicted amino acid sequences where they have been determined for the probe and exon nucleotide sequences. These are set out as PEPTIDE SEQ ID NOS.:. The 5 peptide sequences for a given exon are predicted as follows: Since each chip exon is a consensus sequence drawn from predictions from various exon finding programs (i.e. Grail, GeneFinder and GenScan), the multiple initial ORFs are first determined in a uniform way according to each 10 prediction. In particular, the reading frame for predicting the first amino acid in the peptide sequence always starts with the first base of any codon and ends with the last base of non-termination codon. Next, for each strand of the exon, initial ORFs are merged into one or more final ORFs in an exhaustive process based on the following criteria: 1) the merging ORFs must be overlapping, and 2) the merging ORFs must be in the same frame.

The Sequence Listing, which is a superset of all of the data presented in Table 4, further includes, for 20 each probe, the most similar hit, with accession number and BLAST E value, from the each of the three queried databases.

Table 4 further lists, for each probe, a portion of the descriptor for the top hit ("Top Hit Descriptor") as 25 provided in the sequence database. For those ORFs that are similar in sequence, but nonidentical to known sequences (e.q., those with BLAST E values between about 1e-05 and 1e-100), the descriptor reveals the likely function of the protein encoded by the probe's ORF.

Using BLAST E value cutoffs of 1e-05 (i.e., 1 x 10^{-5}) and 1e-100 (i.e., 1×10^{-100}) as evidence of similarity to sequences known to be expressed is of course arbitrary: in Example 2, supra, a BLAST E value of 1e-30 was used as the boundary when only two classes were to be defined for 35 analysis (unknown, >1e-30; known <1e-30) (see also FIG. 8).

Furthermore, even when the "Most Similar (Top) Hit BLAST E Value" is low, e.g., less than about 1e-100 — which is probative evidence that the query sequence has previously been shown to be expressed — the top hit is highly unlikely exactly to match the probe sequence.

First, such expression entries typically will not have the intronic and/or intergenic sequence present within the single exon probes listed in the Table. Second, even the ORF itself is unlikely in such cases to be present identically in the databases, since most of the EST and mRNA clones in existing databases include multiple exons, without any indication of the location of exon boundaries.

As noted, the data presented in Table 4 represent a proper subset of the data present within the attached

15 sequence listing. For each amplicon probe (SEQ ID NOs.: 1 - 12,673) and probe exon (SEQ ID NOs.: 12,674 - 25,129, respectively), the sequence listing further provides, through iterated annotation fields <220> and <223>:

- (a) the accession number of the BAC from which the sequence was derived ("MAP TO"), thus providing a link to the chromosomal map location and other information about the genomic milieu of the probe sequence;
- (b) the most similar sequence provided by BLAST query of the EST database, with accession number and BLAST 25 E value for the "hit";
 - (c) the most similar sequence provided by BLAST query of the GenBank NR database, with accession number and BLAST E value for the "hit"; and
- (d) the most similar sequence provided by BLASTX 30 query of the SWISSPROT database, with accession number and BLAST E value for the "hit".

EXAMPLE 5

35 Genome-Derived Single Exon Probes Useful For Measuring

Expression of Genes in Human Fetal liver

Table 4 (526 pages) presents expression, homology, and functional information for the genome-derived single exon

5 probes that are expressed significantly in human Fetal liver.

CLAIMS

1. A spatially-addressable set of single exon nucleic acid probes for measuring gene expression in a sample derived

- from human Fetal liver comprising a plurality single exon nucleic probes, said probes comprising any one of the nucleotide sequences set out in SEQ ID NOs: 1 12,673 or a complementary sequence, or a portion of such a sequence.
- 10 2. A spatially-addressable set of single exon nucleic acid probes as claimed in claim 1 wherein each of said plurality of probes is separately and addressably amplifiable.
- 3. A spatially-addressable set of single exon nucleic acid probes as claimed in claim 1 wherein each of said plurality of probes is separately and addressably isolatable from said plurality.
- 4. A spatially-addressable set of single exon nucleic acid 20 probes as claimed in any of claims 1 to 3 wherein said probes comprise any one of the nucleotide sequences set out in SEQ ID NOS.: 12,674 - 25,129.
- 5. A spatially-addressable set of single exon nucleic acid probes as claimed in any of claims 1 to 4, wherein each of said plurality of probes is amplifiable using at least one common primer.
- 6. A spatially-addressable set of single exon nucleic acid 30 probes as claimed in any of claims 1 to 5 wherein the set comprises between 50 - 20,000 single exon nucleic acid probes.
- 7. A spatially-addressable set of single exon nucleic acid probes as claimed in any of claims 1 to 6, wherein the

average length of the single exon nucleic acid probes is between 200 and 500 bp.

- 8. A spatially-addressable set of single exon nucleic acid probes as claimed in any of claims 1 to 7, wherein at least 50% of said single exon nucleic acid probes lack prokaryotic and bacteriophage vector sequence.
- 9. A spatially-addressable set of single exon nucleic acid 10 probes as claimed in any of claims 1 to 8, wherein at least 50% of said single-exon nucleic acid probes lack homopolymeric stretches of A or T.
- 10. A spatially-addressable set of single exon nucleic acid 15 probes as claimed in any of claims 1 - 9 characterised in that said set of probes is addressably disposed upon a substrate.
- 11. A spatially-addressable set of single exon nucleic acid 20 probes as claimed in claim 10 wherein said substrate is selected from glass, amorphous silicon, crystalline silicon and plastic.
- 12. A microarray comprising a spatially addressable set of25 single exon nucleic acid probes as claimed in any of claims1 11.
- 13. A single exon nucleic acid probe for measuring human gene expression in a sample derived from human Fetal liver comprising a nucleotide sequence as set out in any of SEQ ID NOs.: 1 12,673 or a complementary sequence or a fragment thereof wherein said probe hybridizes at high stringency to a nucleic acid molecule expressed in the human Fetal liver.

14. A single exon nucleic acid probe as claimed in claim 13 comprising a nucleotide sequence as set out in any of SEQ ID NOs.: 12,674 - 25,129 or a complementary sequence or a fragment thereof.

5

- 15. A single exon nucleic acid probe for measuring human gene expression in a sample derived from human Fetal liver which is a nucleic acid molecule having a sequence encoding a peptide comprising a peptide sequence as set out in any
- of SEQ ID NOs.: 25,130 37,156, or a complementary sequence or a fragment thereof wherein said probe hybridizes at high stringency to a nucleic acid expressed in the human Fetal liver.
- 15 16. A single exon nucleic acid probe as claimed in any one of claims 13 to 15 wherein said single exon nucleic acid probe comprises between 15 and 25 contiguous nucleotides of said SEQ ID NO.
- 20 17. A single exon nucleic acid probe as claimed in any one of claims 13 to 15, wherein said probe is between 3 25 kb in length.
- 18. A single exon nucleic acid probe as claimed in any one 25 of claims 13 - 17, wherein said probe is DNA, RNA or PNA.
 - 19. A single exon nucleic acid probe as claimed in any one of claims 13 18, wherein said probe is detectably labeled.

- 20. A single exon nucleic acid probe as claimed in any one of claims 13 19, wherein said probe lacks prokaryotic and bacteriophage vector sequence.
- 35 21. A single exon nucleic acid probe as claimed in any one

of claims 13 - 20, wherein said probe lacks homopolymeric stretches of A or T.

- 22. A method of measuring gene expression in a sample 5 derived from human Fetal liver, comprising:
 - contacting the microarray of claim 12, with a first collection of detectably labeled nucleic acids, said first collection of nucleic acids derived from mRNA of human Fetal liver; and then
- measuring the label detectably bound to each probe of said microarray.
 - 23. A method of identifying exons in a eukaryotic genome, comprising:
- algorithmically predicting at least one exon from genomic sequence of said eukaryote; and then detecting specific hybridization of detectably labeled nucleic acids to a single exon probe,
- wherein said detectably labeled nucleic acids are derived
 from mRNA from the Fetal liver of said eukaryote, said
 probe is a single exon probe having a fragment identical in
 sequence to, or complementary in sequence to, said
 predicted exon, said probe is included within a microarray
 according to claim 12, and said fragment is selectively
 hybridizable at high stringency.
 - 24. A method of assigning exons to a single gene, comprising:

- identifying a plurality of exons from genomic sequence according to the method of claim 23; and then
 - measuring the expression of each of said exons in a plurality of tissues and/or cell types using hybridization to single exon microarrays having a probe with said exon.

wherein a common pattern of expression of said exons in said plurality of tissues and/or cell types indicates that the exons should be assigned to a single gene.

- 5 25. A nucleic acid sequence as set out in any of SEQ ID NOs: 1 25,129 which encodes a peptide.
 - 26. A peptide encoded by a sequence as set out in any of . SEQ ID Nos: 1 25,129.

10

27. A peptide comprising a sequence as set out in any of SEQ ID Nos: 25,130 - 37,156.

Page 1 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

בווקום ראכון בנסומה באלוים מימי ביי מימי	Top Hit Descriptor																																		
EXOIL FIGURES	Top Hit Database Source																																		
Single	Top Hit Acesslon No.																																		
	Most Similar (Top) Hit BLAST E																																		
	Expression Signal	4.41	8.8	2.9	10.32	2.59	5.03	1.73		9.24	1.21	3.24	4.38	2.04	0.89		1.65	1.22	10.28	9.0	16.0	0.94	1.53	8.4	0.74	0.74	1.3	1,04	0,59		5.95	1.32			5.64
	ORF SEQ ID NO:	25600						_	26919				27360	27465	27735		28311	28578			28787		28310			29397		29979		30071	30187	30198		30418	
	Exen SEQ ID NO:	13108	L_		13940	14248	14270	14353	14375		14519		14786	14890	15169		15832	16101	16170	16220	16319	16618	16861	16935	16955	16955	17016	17537	17580	L	17762	17777	Ш	1 I	18244
	Probe SEQ ID NO:	475	922	1083	1345	1658	1678	1763	1785	1792	1935	2021	2210	2318	2607	2807	3220	3496	3566	3617	3718	4020	4275	4348	4368	4368	4430	4962	2005	5054	5197	5212	5482	5462	5615

Page 2 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor															والمراجع المراجع المرا																			
Top Hit Database Source																																		
Top Hit Acession No.																																		
Most Similar (Top) Hit BLAST E Vætue																																		
Expression Signal	9.03	4.85	0.84	3.16	1.41	1.65	1.26	-	1	1.13	1.13	1.4	4.1	1.65	1.45	0.57	0.57	4.84	0.78	1.19	1.03	0.48	0.48	0.65		3.06	2.46	2.99	2.73	1.87	1.87	2.59	2.19	1.6
ORF SEQ ID NO:			31257	31262	31552			32067		32569	32560	32831	32832	33451			34258	34931	35155			35702	35703		35816		36268		36827	36043				30914
SEQ ID	18408	18244	18532	18537	24759	18810	19146	19264	19264	19711	19711	19965	19965	20547	20962	21333	21333	21979	22180	22294		22709	22709			23099		23425	<u> </u>	23034	1	1		I J
Probe SEQ ID NO:	5783	5859	5910	5915	6173	9200	6548	8999	8999	7179	7179	7441	7441	8005	8422	8794	8794	9453	9681	9426	9836	10214	10214	10328	10326	10563	10725	10906	11238	11338	11336	11374	12117	12439

Page 3 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Homo sepiens LSS gene, partial, exons 15, 16, 17 and 18	Haemophilus influenzae Rd section 31 of 163 of the complete genome	Sulfolobus soffataricus 281 kb genomic DNA fragment, strain P2	Sulfolobus soffataricus 281 kb genomic DNA fragment, strain P2	Gallus gallus omithine transcarbamylase (OTC) gene, exon 1	Gallus gallus ornithine transcarbamylase (OTC) gene, exon 1	Mus musculus Najb3 gene, exon 1; neuronal apoptosis inhibitory protein 1 (Najp1) and general transcription factor IIH polypeptide 2 (Gtf2h2) genes, complete cds	Mus musculus Nalp3 gene, exon 1; neuronal apoptosis inhibitory protein 1 (Naip1) and general transcription factor IIH polypoptide 2 (Gtf2h2) genes, complete cds	Dengue virus type 3 membrane protein (prM/M)/envelope glycoprotein (E) polyprotein mRNA, partal cds	Dengue virus type 3 membrane protein (pr/M/M)/envelope glycoprotein (E) polyprotein mRNA, perital cds	Mus musculus AT3 gene for antithrombin, complete cds	Homo sapiens ectodysplasin-A receptor protein (EDAR) gene, exons 2, 3, and 4	IMMEDIATE-EARLY PROTEIN 1 (IE1) (IMMEDIATE-EARLY PHOSPHOPROTEIN PP89)	Leuciscus cephalus orientalis cytochroma b (cyt b) gene, partial ods; mitochondrial gene for mitochondrial product	Leuciscus cephalus orientalis cytochrome b (cyt.b) gene, partial ods; mitochondrial gene for mitochondrial products	RHODOPSIN	801851038R1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:3934592 3'	Cynops pyrrhogaster CpTbx3 premature mRNA, partial cds	Cynops pyrrhogaster CpTbx3 premature mRNA, partial cds	Homo sapiens insulin receptor substrate 1 (IRS1) mRNA	Zea mays mRNA for legumain-like protease (see2a)	BREFELDIN A RESISTANCE PROTEIN	African swine fever virus NP1450L gene encoding RNA polymerase largest subunit	Thermoplasma acidophitum complete genome; segment 3/5	THROMBOSPONDIN 1 PRECURSOR	THROMBOSPONDIN 1 PRECURSOR	602128876F1 NIH_MGC_56 Homo sapiens cDNA clone IMAGE:4285508 5'	HISTIDINE-RICH GLYCOPROTEIN PRECURSOR
Top Hit Database Source	H	H	NT S	NT S		NT	NT fe	N 5 35	_ O	a tn	N TN		SWISSPROT	J N	N	ISSPROT	EST_HUMAN 6	Į.			Z	Г	Ā	Н	SWISSPROT			SWISSPROT
Top Hit Acession No.	9.9E+00 AJ239028.1					9.6E+00 AF065630.1	9.6E+00 AF242432.1	9.6E+00 AF242432.1			8.4E+00 AB043785.1		P11210	9.1E+00 AF095609.1			8.9E+00 BE971806.1	8.7E+00 AB019788.1	8.7E+00 AB019788.1	5031804 NT	8.1E+00 AJ131719.1		-	 -	P35441	P35441	7.4E+00 BF700517.1	P04929
Most Similar (Top) Hit BLAST E Value	9.9E+00	9.8E+00 U32716.1	9.8E+00 Y18930.1	9.8E+00 Y18930.1	9.6E+00	9.6E+00	9.6E+00	9.6E+00	9.4E+00 L11433.1	B.4E+00 L11433.1	9.4E+00	9.3E+00	9.3E+00 P11210	9.1E+00	0 4 11 + 00	9.0E+00	8.9E+00	8.7E+00	8.7E+00	8.4E+00	8.1E+00	8.0E+00 P41820	7.6E+00	7.5E+00	7.5E+00 P35441	7.5E+00 P35441	7.4E+00	7.4E+00 P04929
Expression Signal	14.37	1.85	0.47	0.47	0.8	8.0	1.22	1.22	1.14	1.14	3.18	0.99	3.48	2.82	2.82	6.0	6.12	1.9	1.9	1.68	3.8	2.47	0.78	1.95	1.54	1.54	3.35	2.63
ORF SEQ ID NO:	31583	33400	35128	35129		32484	35808		27814	27815	28040		34390	30543	30544	L	31564	31907	31908		33217				33764			34147
Exan SEQ ID NO:	18813	20490	l			19645	22813		15247	15247	15588	20584		18134	18134	L	L		19117	13099	20315	23581	20633		20843	20843	ll	21227
Probe SEQ ID NO:	6203	7948	898	9658	7073	7073	10319	10319	2889	2883	956Z	8042	8933	2500	7500	9351	6186	6517	6517	465	9376	11048	8092	7384	8302	8302	5968	8888

Page 4 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

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Top Hit Descriptor		HISTIDINE-RICH GLYCOPROTEIN PRECURSOR	Lycopersicon esculentum Mill. GTPase (SAR2) mRNA, complete cds	Lycopersicon esculentum Mill. GTPase (SAR2) mRNA, complete cds	RC0-HT0613-200300-031-a07 HT0613 Homo sapiens cDNA	ZINC-FINGER PROTEIN 1 (ZINC-FINGER HOMEODOMAIN PROTEIN 1)	ZINC-FINGER PROTEIN 1 (ZINC-FINGER HOMEODOMAIN PROTEIN 1)	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 91	HYPOTHETICAL 17.3 KDA PROTEIN IN MRDA-PHPB INTERGENIC REGION	ARGININE KINASE (AK)	WD-40 REPEAT PROTEIN MSI3	60S RIBOSOMAL PROTEIN L4 (L2)	DNA MISMATCH REPAIR PROTEIN MUTS	2807c11.r1 Soares melanocyte 2NbHM Homo sapiens cDNA clone IMAGE:291860 5'	za07c11.r1 Sceres melanocyte 2NbHM Homo sapiens cDNA clone IMAGE:291880 5'	OUTER CAPSID PROTEIN VP4 (HEMAGGLUTININ) (OUTER LAYER PROTEIN VP4) [CONTAINS:	OUTER CAPSID PROTEINS VP5 AND VP8]	HYPOTHETICAL 157.0 KDA PROTEIN C38C10.5 IN CHROMOSOME III	CATECHOL-O-METHYLTRANSFERASE, SOLUBLE FORM (S-COMT)	URIDYLATE KINASE (UK) (URIDINE MONOPHOSPHATE KINASE) (UMP KINASE)	URIDYLATE KINASE (UK) (URIDINE MONOPHOSPHATE KINASE) (UMP KINASE)	PROBABLE CATION-TRANSPORTING ATPASE C8C3.05C	ENV POLYPROTEIN (CONTAINS: COAT PROTEIN GP52, COAT PROTEIN GP36)	601678435F1 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3960969 5'	Schizophyllum commune unknown mRNA	Mus musculus mannosidase 2, alpha B1 (Man2b1), mRNA	601468031F1 NIH_MGC_67 Homo sapiens cDNA clone IMAGE:3871303 5	Pyrococcus harikashii OT3 genamic DNA, 1188001-1485000 m. position (8/7)	Deinococcus radiodurans R1 section 1 of 2 of the complete chromosome 2	Deinococcus rediodurans R1 section 1 of 2 of the complete chromosome 2	Mus musculus mixed lineage kinase 3 (MIK3) and two pore domain K+ channel subunit (Kcnk8) genes,	complete cds	Homo sapiens DESC1 protein (DESC1); mRNA	Mus musculus immunoglobulin scavenger receptor IgSR mRNA, complete cds	Mus musculus immunoglobulin scavenger receptor IgSR mRNA, complete cds
Top Hit Database Source		SWISSPROT	L	LN	EST_HUMAN	SWISSPROT	SWISSPROT	NT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	EST_HUMAN	EST_HUMAN		SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	EST_HUMAN	NT	NT	EST_HUMAN	LN TN	N	N		Ā	Z	ΙΝ	TN
Top Hit Acession No.		04929	12051.1	.12051.1	7.2E+00 BE179090.1	28166	28166	7.1E+00 AL181595.2	205850				544834	W03412.1	W03412.1		>36307	203570				210309	P03374	6.5E+00 BE866001.1	6.2E+00 AY010901.1	6754621 NT	6.0E+00 BE780163.1	6.0E+00 AP000006.1	AE001862.1	6.0E+00 AE001862.1		5.9E+00 AF155142.1	7681557 NT	5.7E+00]AF302046.1	5.7E+00 AF302046.1
<u>a</u> = 111	Value	7.4E+00 P04929	7.2E+00	7.2E+00 L12051.1	7.2E+00	7.1E+00 P28168	7.1E+00 P28166	7.1E+00	7.1E+00 P05850	7.0E+00 P48610	7.0E+00	6.9E+00 P35679	6.9E+00 P44834	6.8E+00 W03412.1	6.8E+00 W03412.1		6.8E+00 P36307	6.8E+00 Q03570	6.6E+00 Q99028	00+39:9	6.6E+00 Q9ZE07	6.6E+00 Q10309	6.5E+00 P03374	00+3€-90	6.2€+00	6.2E+00	6.0E+00	6.0E+00	00+30.9	6.0E+00		5.9E+00	5.8E+00	5.7E+00	5.7E+00
Expression Signal		2.63	3.19	3.19	0.7	1.22	1.22	7.96	3.2	3.35	1.87	4.08	1.2	1.38	1.38		1.13	3.85	0.69	1.89	1.89	2.13	7.21	0.49	1.11	,0.53	1.34	0.46	9.0	9.0		6.67	1.18	0.67	19:0
ORF SEQ ID NO:		34148	28099	28100	32507	32585	32586		36848	35367	36679	33684		33283	33294			35595		35465	35466		34584	35695	35127			35205	35896	35897		32042		32601	32602
Excan SEQ ID				15622	19668	19734	19734		23791		23637			20391	20391		21597	22605	18122	22481	22481	23522	21644	10/22		22949			22901		l				19746
Probe SEO ID		8898	3006	3008	7097	7203	7203	9218	11263	8892	11129	8225	10253	7849	7849		906	10110	5488	9866	9866	11008	9108	10206	9657	10455	7102	9730	10407	10407		6843	3576	7215	7215

Page 5 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 6 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 7 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

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Single Exoll Probes Expressed in Total	Top Hit Descriptor	CYCLIN-DEPENDENT KINASE INHIBITOR 1B (CYCLIN-DEPENDENT KINASE INHIBITOR P27) (P27KIP1)	3-OXOACYL-[ACYL-CARRIER-PROTEIN] SYNTHASE III (BETA-KETOACYL-ACP SYNTHASE III) (KAS II)	HYPOTHETICAL PROTEIN HVLF1	601507510F1 NIH_MGC_71 Homo saplens cDNA clone IMAGE:3909051 5'	GLC7-INTERACTING PROTEIN 1	SUCRASE-ISOMALTASE, INTESTINAL [CONTAINS: SUCRASE ; ISOMALTASE]	SUCRASE-ISOMALTASE, INTESTINAL (CONTAINS: SUCRASE; ISOMALTASE)	SUCRASE-ISOMALTASE, INTESTINAL [CONTAINS: SUCRASE; ISOMALTASE]	SUCRASE-ISOMALTASE, INTESTINAL [CONTAINS: SUCRASE; ISOMALTASE]	CELL DIVISION PROTEIN FTSY HOMOLOG	Ureaplasma urealyticum section 33 of 59 of the complete genome	URICASE (URATE OXIDASE)	URICASE (URATE OXIDASE)	GENOME POLYPROTEIN [CONTAINS: CAPSID PROTEIN C (CORE PROTEIN); MATRIX PROTEIN (ENVELOPE GLYCOPROTEIN M); MAJOR ENVELOPE PROTEIN E; NONSTRUCTURAL PROTEINS NS1, NS2A, NS2B, NS4 AND NS4B; HELICASE (NS3); RNA-DIRECTED RNA POLYMERASE (NS3)]	GENOME POLYPROTEIN (CONTAINS: CAPSID PROTEIN C (CORE PROTEIN); MATRIX PROTEIN	(ENVELOPE GLYCOPROTEIN M); MAJOR ENVELOPE PROTEIN E; NONSTRUCTURAL PROTEINS INS1, NS28, NS48, AND NS48; HELICASE (NS3); RNA-DIRECTED RNA POLYMERASE (NS5)]	N.tabacum chilinase gene 50 for class I chiltinase C	Mus musculus seminal vesicle secretory protein 99 (MSVSP99) gene, promoter region	MR0-BN0070-300500-028-h05 BN0070 Homo sapiens cDNA	MR0-BN0070-300500-028-h05 BN0070 Homo capiens cDNA	Dictyostellum discoldeum non-LTR retrotransposon TRE5-B, polyprotein (gag) and group-specific antigen (pol) genes, complete cds	Human hereditary haemochromatosis region, histone 2A-like protein gene, hereditary haemochromatosis (HLA-H) gene, RoRet gene, and sodium phosphate transporter (NPT3) gene, complete cds	HYPOTHETICAL TRANSCRIPTIONAL REGULATOR IN AIDB-RPSF INTERGENIC REGION	Human MHC class II lymphocyte antigen (DPw4-beta-1) gene, exon 2
EXOII FIODES	Top Hit Database Source	SWISSPROT	SWISSPROT	SWISSPROT	EST_HUMAN	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	LΝ	SWISSPROT	SWISSPROT	SWISSPROT		SWISSPROT	ΙZ	N	EST_HUMAN	EST_HUMAN	LN.	LZ	SWISSPROT	NT
aifilic	Top Hit Acession No.	P46414	084242	P09716	4.1E+00 BE885880.1	P38229	062653	062653	062653				4.0E+00 Q00511	Q00511	P07564		P07564	3.9E+00 X64518.1	3.9E+00 AF055468.1	_		3.9E+00 AF268209.1	3.9E+00 U91328.1	3.9E+00 P39299	
Ì	Most Similar (Top) Hit BLAST E Value	4.1E+00 P46414	4.1E+00 084242	4.1E+00 P09716	4.1E+00	4.0E+00 P38229	4.0E+00 O62653	4.0E+00 O62653	4.0E+00 O62653	4.0E+00	4.0E+00 O33010	4.0E+00	4.0E+00	4.0E+00 Q00511	4.0E+00 P07564		4.0E+00 P07564	3.9€+00	3.9E+00	3.9E+00	3.9E+00	3.9E+00	3.9E+00	3.9E+00	3.9E+00
	Expression Signal	0.5	0.62	2.97	13.84	0.82	0.74	0.74	0.95	0.95	1.34	9.0	0.49	0.49	3.99		3.99	4.79	0.74	3.08	3.08	0.71	0.72	4.12	60.9
	ORF SEQ ID NO:		36003				32336	32337		32337	32625	35560	35647	35648	2669£		36938			31186			32198	732357	
	Exen SEQ ID NO:	22703	22993	1	23372	L	19515	19515	19515	19515	19769	22565	22653	22653			23874	ł	16998	18463	18463	İ	19383		19923
	Probe SEQ ID NO:	10208	10499	10765	10851	3599	2850	2650	7017	7017	7240	10070	10158	10158	11423		11423	3550	4413	5839	5839	6746	6792	6955	7398

Page 8 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	Exen NO: 23307 23007 23007 23007 23007 23007 23001 24158 13246 11492 21126 211		Signa	8 S C E E C C C C C C C C C C C C C C C C	Top Hit Acession No. No. No. No. Acese65.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 718000.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1 71800.1	▎▝▗▗░▗▕▗▎▗░▗░░▄░▕▕▕▗▕▗▐▄░▀░▕▀▎▜▀▞▀▍▎▘ ▗░▗░	Xleavis mRNA for M4 muscarinic receptor Alexon septens NF2 gene Arrival Septens NF2 gene METALLOPROTEINASE HINBITOR TO PRECURSOR (HUMAN); Helicobacter pykol, strain J89 section 123 of 132 of the complete genome HYPOTHETICAL PROTEIN MJ0385 HUMSUDYT35 Human brain cDNA Homo septens cDNA clone 148 Stratucobase oralis partial strain gene for xambline phosphoribosyltransfrease, strain NCTC7884 Arabidopsis theliana DNA chromosome, segment 3/5 Homo sapiens glucokinese (hexokinese 4, maturity onset diabetes of the young 2) (GCK), nuclear gene encoding mitochordida protein, mRNA Mus musculus terminin beta 2 gene, exons 17-33, and complete cds 602120551F1 NIH_MGC_56 Homo sapiens cDNA clone IMAGE:427748 5 602120551F1 NIH_MGC_56 Homo sapiens cDNA clone IMAGE:4277748 5 Gallus gallus mRNA for hypode-Inducible factor-1 alpha, complete cds Arebidopsis theliane DNA chromosome 4, contig fragment No. 29 HUM000TB08 Liver HepG2 cell line. Homo sapiens cDNA clone tb08 HUM000TB08 Liver HepG2 cell line. Homo sapiens cDNA clone tb08 Fesudormonias aeruginosa PAO1, section 8 of 529 of the complete genome Escherichia coli glycerophosphete deflydrogenses (gIPD) gene, partial cds; and the translation start site has been verified (gipC), and repressor protein (gipR) gene, partial cds complete cds Chytosporidium felix heat shock protein 70 (HSP70) gene, partial cds Gorpusperidium felix heat shock protein (capC) gene, partial cds Gorpusperidium felix heat and the mon sapiens cDNA clone INAGE:34400 5; Borrelia burgdorfer (strain 25015) outer surface protein (capC) gene, partial cds Gorpusperidium felix heat and the homo sapiens cDNA clone INAGE:34400 5;
6360	18964	31742	0.92		R19745.1 P24557	EST HUMAN SWISSPROT	yg40c08.r1 Soares infent brain 1NIB Homo sapiens cDNA clone IMAGE:34940 5' THROMBOXANE-A SYNTHASE (TXA SYNTHASE) (TXS)
8962 8962 9414	1 1	34421				EST_HUMAN EST_HUMAN NT	298504.s1 Stratagene HeLa cell s3 837216 Homo sapiens cDNA clone IMAGE:627055 3' similar to contains Alu repetitive element. 2986b04.s1 Stratagene HeLa cell s3 937216 Homo sapiens cDNA clone IMAGE:627055 3' similar to contains Alu repetitive element; contains element MSR1 repetitive element. Arabidopsis thaliana DNA chromosome 4, contig fragment No. 53

Page 9 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 10 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Bacillus alcalophilus pectate lyase (pelE) gene, complete cds	TYPE I IODOTHYRONINE DEIODINASE (TYPE-1 5DEIODINASE) (DIOI) (TYPE 1 DI) (5DI)	TYPE I IODOTHYRONINE DEIODINASE (TYPE-I S'DEIODINASE) (DIOI) (TYPE 1 DI) (5DI)	GLUTAMATE (NMDA) RECEPTOR SUBUNIT EPSILON 3 PRECURSOR (N-METHYL D-ASPARTATE DECEPTOR SUBTAGE CONVINCION SUBT	ACCELLOR SUBTIFICACIÓN (MACA) (MADANAC)	COLLAGEN ALPHA 2(1) CHAIN PRECURSOR	Chlorella vulgaris chloroplast, complete genome	HYPOTHETICAL 56.3 KD PROTEIN F52C9.5 IN CHROMOSOME III	DEOXYHYPUSINE SYNTHASE (DHS)	GENOME POLYPROTEIN (CONTAINS: CAPSID PROTEIN C (CORE PROTEIN); MATRIX PROTEIN	(ENVELOPE PROTEIN M), MAJOR ENVELOPE PROTEIN E; NONS ROCTORAL PROTEINS NST, NS2A, NS2B, NS4A AND NS4B, HELICASE (NS3); RNA-DIRECTED RNA POLYMERASE (NS5))	retinoic acid nuclear receptor isoform beta 2 [mice, embryonal carcinoma cell line, PCC7-MZ1, mRNA, 2971	(tu	Brassica rapa pollen coat protein homolog (BAN103) gene, complete cds	Saureus genes encoding Sau96! DNA methyltransferase and Sau96! restriction endonuclease	Corynebacterium glutamicum thrC gene for threonine synthase (EC 4.2.99.2)	Corynebacterium glutamicum thrC gene for threonine synthase (EC 4.2.99.2)	CYR61 PROTEIN PRECURSOR (3CH61)	ENDOTHELIAL CELL MULTIMERIN PRECURSOR	B.napus DNA for myrosinase	S-ADENOSYLMETHIONINE SYNTHETASE (METHIONINE ADENOSYLTRANSFERASE) (ADOMET	CDC16 PROTEIN HOMOLOG	RETINAL GUANYLYL CYCLASE 2 PRECURSOR (GUANYLATE CYCLASE 2F, RETINAL) (RETGC-2)	(ROD OUTER SEGMENT MEMBRANE GUANYLATE CYCLASE 2) (ROS-GC2) (GUANYLATE CYCLASE	F) (GC-F)	RETINAL GUANYLYL CYCLASE 2 PRECURSOR (GUANYLATE CYCLASE 2F, RETINAL) (RETGC-2)	(ROD OUTER SEGMENT MEMBRANE GUANYLATE CYCLASE 2) (ROS-GC2) (GUANYLATE CYCLASE	F) (GC-F)	Chlamydophila pneumoniae AR39, section 53 of 94 of the complete genome	Bonapartia pedaliota mitochondrial DNA for 16S ribosomal RNA	F.pringlei gdcsPA gene for P-protein of the glycine cleavage system	
Top Hit Database Source	Ł	SWISSPROT	SWISSPROT	Fordonisio	SWISSPROI	SWISSPROT	L	SWISSPROT	SWISSPROT		SWISSPROT		Ā	IN	٦	F	L	SWISSPROT	SWISSPROT	Ę	100000000000000000000000000000000000000	SWISSPROT			SWISSPROT			SWISSPROT	L	NT	NT	
Top Hit Acession No.	AF303225.1	P49894	P49894	1,077	014937	Q01149	7524759 NT	Q10125	P49365		P33515		S56660.1	17666.1	X53096.1	X56037.1	X56037.1	P18406	Q13201	X67838.1		016181			P51842			P51842	AE002225.2	AB026033.1	236879.1	,,,,,
Most Similar (Top) Hit BLAST E Value	3.1E+00	3.1E+00	3.1E+00			3.1E+00			3.1E+00		3.1E+00		3.1E+00	3.1E+00	3.0E+00.	3.0E+00	3.0E+00	3.0E+00		3.0E+00		305-100			3.0E+00			3.0E+00	2.9E+00	2.9E+00	2.9E+00 Z36879.1	100.300
Expression Signal	1.09	4.27	4.27	į	27.5	0.52	0.75	0.56	4.7		2.91		7.48	1.38	1.68	0.72	0.72	10.44	0.77	1.33		1.62			7.04		_	7.04	2.32	0.68	3.74	166 1
ORF SEQ ID NO:		33995				34726			35726						30588	32073	32074					36075			36426				27207		32309	
Exan SEQ ID NO:	20220	ı	21077	3	П		ı	22398	22734		23036	ı	23809	24819	18173	19269	19269	19740	19776	21377	l	23064	1		23409				14636	18833	19487	40700
Probe SEQ ID NO:	7711	8538	8538		3 5	9249	8 5	9899	10239		11338		11355	12490	5541	6873	6873	7209	7247	8838		10527			10888			10888	2055	6224	6869	7282

Page 11 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

SEQ Expression (Top) Hit Acession Signal BLASTE No. Source Source	32845 4.37 2.9E+00 014514 SWISSPROT BRAIN-SPECIFIC ANGIGGENESIS INHIBITOR 1 PRECURSOR	6.04 2.9E+00 P46589 SWISSPROT	33280 0.67 2.9E-400 P05844 SWISSPROT NONSTRUCTURAL PROTEIN VP4; MINOR STRUCTURAL PROTEIN VP2; NONSTRUCTURAL PROTEIN VP3]	0.67	0.89 2.9E+00 BF344171.1 EST_HUMAN	4.87 2.8E+00 AF186398.1 NT	3.45 2.8E+00 AL161552.2 NT Arebidopsis thaliana DNA chromosome 4, contig fregment No. 52	4.88 2.8E+00 8393724[NT	0.57 2.8E+00 BE565182.1 EST_HUMAN	1.68 2.8E+00] 8393724 NT	25394 9.31 2.7E+00 6676306 NT Mus musculus per-höxemer repeat gene 3 (Phx3), mRNA	25395 9.31 2.7E+00 6678306 NT Mus musculus per-hexamer repeat gene 3 (Phor3), mRNA	31073 1.2 2.7E+00 L14005.1 IVT Homo sapiens apoA polymorphism Kringle IV gene, exons 1 and 2	U15947.1 NT	1.69 2.7E+00 AL116459.1 NT Botryts cinerea strain T4 cDNA library under conditions of nitrogen deprivation	0.63 2.7E+00 AW088191.1 EST_HUMAN	1.48 2.7E+00 BE063527.1 EST_HUMAN CM0-BT0281-031199-087-h04 BT0281 Homo eapliens cDNA		31068 1.94 2.6E+00 6755601 NT Mus musculus SRY-box containing gene 13 (Sox13), mRNA	1.94	2.42 2.6E+00[Y17062.1 INT Mycobacterium fortuitum furA II gene	5.98 2.6E+00 AF235502.1 NT Mus musculus SH2-containing inositel 5-phosphatase (Ship) gene, exons 16 through 27, and complete cds	1.08 2.6E+00 AJ132180.1	1.08 2.6E+00[AJ132180.1 NT	2.6E+00 AL161540.2 NT	1.51 2.8E+00 9055193 NT	36443 1.69 2.8E+00 AF143875.1 INT [Hantavirus Z10 segment M G1/G2 glycoprotein (Z10) gene, complete cds	11419220 NT	2.29 2.5E+00 AJ271844.1 NT	2.29
							3.45							9.0	1.68		1.48				2.42	5.98						2.76		
ORF SEQ ID NO:	32845	32868	33260	33261			21	32740					31073	63	91	33191	g		32 31068	31069	12		15 33447				36443	92		
Exon SEQ ID NO:	19790	20001	7809 20352	7809 20352		1504 14096	1875 14267	19874		10569 19874	12911		18368	3 20629	3 21436	20292	22888	17362	18362	18382	18612	20198	3 20545				7 23428	24986	14105	14105
Probe SEQ ID	7262	7479				• •	40	7348	9531	3	251	251	5740	8088	8898	9353	10394	4781	5738	5736	5992	7689	8003	8003	9256	10257	10907	12390	1513	1513

Page 12 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

					,		
Probe SEQ ID NQ:	Exen SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Detabase Source	Top Hit Descriptor
5981	18601	31334	1.71	2.5E +00	P13485	SWISSPROT	TEICHOIC ACID BIOSYNTHESIS PROTEIN F
5981	18601	31335	1.71	2.5E+00	P13485	SWISSPROT	TEICHOIC ACID BIOSYNTHESIS PROTEIN F
6586	18601	31334	1.39	2.5E+00 P13485		SWISSPROT	TEICHOIC ACID BIOSYNTHESIS PROTEIN F
6586	18601	31335	1.39	2.5E+00 P13485		SWISSPROT	TEICHOIC ACID BIOSYNTHESIS PROTEIN F
6828	19418		62'0	2.5E+00 D30052.1		NT	Vibrio chalerae ctxA gene and ctxB gene for chalera toxins, complete cds
7736	20244	33135	1.05	2.5E+00	0 AW949158.1	EST_HUMAN	QV4-FT0005-110500-205-g07 FT0005 Homo sapiens cDNA
9032	21569			2.5E+00		NT	Rice DNA for aidclase C-1, complete cds
926		35247	88.0	2.5E+00	Γ	EST_HUMAN	601175779F1 NIH_MGC_17 Hamo sapiens cDNA clone IMAGE:3531090 5'
11724	24131		1.66	2.5€+00	2.5E+00 AF289665.1	NT	Mus musculus EIF4H gene, partial cds; LIMK1 gene, complete cds; and ELN gene, partial cds
3047	15663	28144	6.0		-	LN	Chicken alpha-3 collagen type VI mRNA, 3' end
5033	17607				4503352 NT	L	Homo sapiens double C2-like domains, alpha (DOC2A) mRNA
6161	18774	31536				SWISSPROT	VITELLOGENIN I PRECURSOR (YOLK PROTEIN 1)
8082	20624	33536				SWISSPROT	CD27L RECEPTOR PRECURSOR (T-CELL ACTIVATION ANTIGEN CD27) (T14)
8082	20624	33537	1.99	2.4E+00 P26842		SWISSPROT	CD27L RECEPTOR PRECURSOR (T-CELL ACTIVATION ANTIGEN CD27) (T14)
8153			2.33	2.4E+00		NT	Helloobacter pylori, strain J99 section 47 of 132 of the complete genome
8585	21124			2.4E+00	2.4E+00 AW875126.1	EST_HUMAN	RC2-PT0004-031299-011-d05 PT0004 Homo sapiens cDNA
8762	21301	34222				SWISSPROT	ENDOCHITINASE B PRECURSOR (CHN-B)
9951	22448		2.59		P13673	SWISSPROT	SKIN GRANULE PROTEIN PRECURSOR
9951	22446				P13673	SWISSPROT	SKIN GRANULE PROTEIN PRECURSOR
10017		35504		'	2.4E+00 X92511.1	LN	H.saplens CTGF gene and promoter region
10141	L		7.38	2.4E+00 P09099	P09089	SWISSPROT	XYLULOSE KINASE (XYLULOKINASE)
10225	22720	35710			2.4E+00 BE326702.1	EST_HUMAN	hr63f08.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:3133187 3'
10225	j				2.4E+00 BE326702.1	EST_HUMAN	hr63f06.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:3133187 3'
10483		35986			051481	SWISSPROT	DENITRIFICATION REGULATORY PROTEIN NIRQ
10958	23473	36498	1.69		2.4E+00 Y14079.1	LN.	Bacillus subtilis chromosomal DNA, region 75 degrees: glpPFKD operon and downstream
11237	23768	36826	2.27		2.4E+00 AF158652.2	Z	Fragaria x ananassa cytosolic ascorbate percoadase (ApxSC) gene, ApxSC-c allete, complete cds
1296					2.3E+00 Z46724.1	NT.	G.domesticus artificial single chain antibody gene (L3)
4199		-	1.65		2.3E+00 AJ401081.1	ΙN	Bos taurus partial cyto gene for cytochrome b
0009	18620		0.91		2.3E+00 N86245.1	EST_HUMAN	J7340F Human fetal heart, Lambda ZAP Express Homo sapiens cDNA clone J7340 5' similar to PROLYLCARBOXYPEPTIDASE
7477	19999	32884			6978554 NT	Ę	Rattus norvegicus ATPase, Ca++ transporting, ubiquitous (Atp2a3), mRNA
7593	3 25120	Ì				SWISSPROT	MAJOR CENTROMERE AUTOANTIGEN B (CENTROMERE PROTEIN B) (CENP-B)
7756	3 20264	33159	1.06		┯.	ĽΝ	M.mazei dhak and dhaJ genes homologues coding for DhaK and DhaJ

Page 13 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor Source		ALPHA-(1,3)-FUCOSYLTRANSFERASE (GALACTOSIDE 3-L-FUCOSYLTRANSFERASE) SWISSPROT (FUCOSYLTRANSFERASE 4) (FUCT-IV)	SWISSPROT ANNEXIN VII (SYNEXIN)	EST_HUMAN 602069121F1 NIH_MGC_58 Homo sapiens cDNA clone IMAGE: 4068173 5'	EST_HUMAN 602069121F1 NIH_MGC_58 Homo sapiens cDNA clone IMAGE:4068173 5'	EST_HUMAN 601433673F1 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:3918643 5',	NT Megnaporthe grisea Class IV chitin synthase (chs4) gene, complete cds	NT Rat gene for regucalcin, exon1 (non-coding exon)	NT Rat gene for regucalcin, exon1 (non-coding exon)	SORTILIN-RELATED RECEPTOR PRECURSOR (SORTING PROTEIN-RELATED RECEPTOR CONTAINING LDLR CLASS A REPEATS) (MSORLA.1) (LOW-DENSITY LIPOPROTEIN RECEPTOR RELATIVE WITH 11 LIGAND-BINDING REPEATS) (LDLR RELATIVE WITH 11 LIGAND-BINDING REPEATS)	SWISSPROT BINDING REPEATS) (LR11) (>	SORTILIN-RELATED RECEPTOR PRECURSOR (SORTING PROTEIN-RELATED RECEPTOR	CONTAINING LOLR CLASS A REPEATS) (MSORLA) (SORLA-1) (LOW-DENSITY LIPOPRO (EIN DECEPTOR RELATIVE WITH 11 LICAND-RINDING REPEATS) (LIDENDENTY WITH 11 LICAND-RINDING REPEATS)	SWISSPROT BINDING REPEATS) (LR11) (>	EST_HUMAN RC3-CT0254-300800-022-e06 CT0254 Homo saplens cDNA	EST_HUMAN RC3-CT0254-300800-022-e06 CT0254 Homo sapiens cDNA	EST_HUMAN 600943401T1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:2959777 3'	SWISSPROT MINOR VIRION STRUCTURAL PROTEIN MU-2	SWISSPROT [INSULIN-LIKE GROWTH FACTOR II PRECURSOR (IGF-II) (SOMATOMEDIN A)	EST_HUMAN IN95b02.s1 NCI_CGAP_Co10 Homo sapiens cDNA clone IMAGE:1058379 3'	EST_HUMAN zn97/04.r1 Stratagene fetal retina 937.202 Homo saplens cDNA clone IMAGE:588143 5'		bb17h12.x1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:2983207 3' similar to gb:D45838 Mouse EST_HUMAN mRNA for nuclear pore-targeting-complex component of (MOUSE);	П	EST_HUMAN mRNA for nuclear pore-targeting-complex component of (MOUSE);		SWISSPROT TRANSPOSON TY1 PROTEIN A	'S SOURCE TO THE PROPERTY OF T
Top Hit Acession No.	5835317 NT	0 011127	907076	2.3E+00 BF541987.1	2.3E+00 BF541987.1	D BE895237.1	AF020528.1	2.2E+00 D67071.1	2.2E+00 D67071.1		088307			0 088307	2.2E+00 BE927220.1	2.2E+00 BE927220.1	2.2E+00 BE 250383.1	000335	2.2E+00 P51459	AA594574.1	2.2E+00 AA137027.1	2.2E+00 AA449012.1	2.2E+00 BE301560.1		2.2E+00 BE301560.1	2.2E+00 BE741678.1	Q04706	
Most Similar (Top) Hit BLAST E Value	2.3E+00	2.3E+00	2.3E+00 Q07076	2.3E+00	2.3E+00	2.3E+00	2.2E+00	2.2E+00	2.2E+00	_	2.2E+00 O88307			2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00	2.2E+00		2.2€+00	2.2€+00	2.2E+00	
Expression Signal	0.53	1.79	2	2.92	2.92	7.31	91.07	4.5	4.5		12.27			12.27	0.95	0.95	9.1	4.32	3.04	3.58	6.0	25.23	0.65		0.65	12.17	2.57	
ORF SEQ ID NO:	34505	34572			37120	31020	29143	29432	29433		30591			30592	31373		31593	31880	32107		32747		33494		33495			
SEQ 1D NO:	21575	1	23213	24055	1	24278	16685	16988	16988		18177			18177	18635		18822	19098	19303	18057	19884	20115	20588	ŀ _	20588	21791		
Probe SEQ ID NO:	8038	7606	10681	11812	11612	11950	4089	4403	4403		5545			5545	8018	8018	6212	6495	6709	7037	7358	7602	8048		8046	9265	9488	

Page 14 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

					Single	Exon Propes	Single Exon Probes Expressed in Fetal Liver
Probe E) SEQ ID SE(Exan OF SEQ ID III	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Detabase Source	Top Hit Descriptor
6966	22461	35444	86.	2.2E+00	AI290373.1	EST_HUMAN	qm69b03.x1 Soares_placenta_8b9weeks_2NbHP8tc9W Homo sapiens cDNA clone IMAGE:1893965 3' similar to gb:Y00433 GLUTATHIONE PEROXIDASE (HUMAN);
10008 2	22503	35494	3.7	2.2€+00	BF246782.1	EST_HUMAN	801855591F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4075391 5'
10353 2	22847	35841	2.99	2.2E+00	AF183416.1	NT	Homo sapiens overian granulosa cell 13.0 kDa protein hGR74 homolog mRNA, complete cds
11316 2	23014	36023	4.01	2.2E+00	P07911	SWISSPROT	UROMODULIN PRECURSOR (TAMM-HORSFALL URINARY GLYCOPROTEIN) (THP)
11482 2	23832	37003	4.23	2.2E+00	P10407	SWISSPROT	EARLY E1A 28 KD PROTEIN
595	15419	25699	6.28	2.1E+00	AF132612.2	NT	Mus musculus pre-T cell receptor alpha gene, enhancer region and upstream region
3648 1	16251		0.65	2.1E+00	AW 449366.1	EST_HUMAN	UI-H-BI3-aki-e-08-0-UI.s1 NCI_CGAP_Sub5 Homo sapiens cDNA clone IMAGE:2734550 3
6281 1	18889		0.85	2.1E+00	P75357	SWISSPROT	HYPOTHETICAL PROTEIN MG302 HOMOLOG
1 6889	19633	32471	3.38	2.1E+00	070159	SWISSPROT	ALPHA-2-HS-GLYCOPROTEIN PRECURSOR (FETUIN-A)
	19450	32266	5.13		N29575.1	EST HUMAN	yydart0.s1 Soares metanocyte 2NbHM Homo sapiens cDNA clone IMAGE:270618 3' similar to gb:M55654 TRANSCRIPTION INITIATION FACTOR TFIID (HUMAN);
	20974		2.27	2.1E+00	AU123630.1	EST_HUMAN	AU 123630 NT 2RM2 Homo sapiens cDNA clone NT 2RM 2000671 5'
10454 2	22948		0.58	2.1E+00	Y10284.1	N	H.sapiens TRAF1 gene, putative promoter region
1238	13836	26352	1.3	2.0E+00	AF180527.1	N	Homo saplens p22Dokdel (DOKDEL) mRNA, complete cds
	13836	26353	1.3	2.0E+00	AF180527.1	TN	Homo sapiens p22Dokdel (DOKDEL) mRNA, complete cds
1380	13973	26501	0.92	2.0E+00	AF204927.1	· LN	Oryctolegus cuniculus Na+,K+-ATPase beta 1 subunit mRNA, complete cds
1619	14212		2.89	2.0E+00	P25582	SWISSPROT	PUTATIVE RRNA METHYLTRANSFERASE SPB1
2194	14770	27343	3.69	2.0E+00	778279.1	NT	R.norvegicus mRNA for collagen alpha1 type I
	14770	27344	3.69	2.0E+00	278279.1	NT	R. norvegicus mRNA for collagen alpha1 type I
	16767	29215	1.9	2.0E+00	AW664496.1	EST HUMAN	hi13c05.x1 NCI_CGAP_GU1 Homo sapiens cDNA clone IMAGE:2972168 3' similar to gb:X01677 GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE, LIVER (HUMAN);
	16767	29216	1.0	<u> </u>	AW664496.1	EST_HUMAN	ht13c05.x1 NCI_CGAP_GU1 Homo sapiens cDNA done IMAGE:2972168 3' similer to gb:X01677 GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE, LIVER (HUMAN);
7552 2	20071		72.0	2.0E+00	P07566	SWISSPROT	STRUCTURAL POLYPROTEIN [CONTAINS: NUCLEOCAPSID PROTEIN C; MEMBRANE GLYCOPROTEINS E1 AND E2]
	20509	33415	3.56	2.0E+00	AB008676.1	۲	Escherichia coli 0157 DNA, map position at 46 min., complete cds
7967	20509	33416	3.56	2.0E+00	AB008676.1	TN	Escherichia coli 0157 DNA, map position at 46 min., complete cds
7967	20509	33417	3.56	2.0E+00	l	TN	Escherichia coli 0157 DNA, map position at 46 min., complete cds
8853	21392	34314	3.62		F31500.1	EST_HUMAN	HSPD22703 HM3 Homo sapiens cDNA clone s4000117808
12295	24946	30622	7.77	2.0E+00	5834843 NT	LN	Gallus gallus mitochondrion, complete genome
	18409	31124				LΝ	Mus musculus inositol 1,4,5-triphosphate receptor 1 (ltpr1), mRNA
	18409	31125	6.89				Mus musculus inosital 1,4,5-triphosphate receptor 1 (Itpr1), mRNA
6249	18858	31630	1.2	1.9E+00	BE969695.1	EST_HUMAN	601679636F1 NIH_MGC_78 Homo sapiens cDNA clone IMAGE:3949881 5

Page 15 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

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Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Тар Hit Descriptar
6760	19353		1.02	1	AW845689.1	EST_HUMAN	MR0-CT0063-071099-002-g02 CT0063 Homo sepiens cDNA
6845	19435		2.31	1.9E+00	Q63627	SWISSPROT	CTD-BINDING SR-LIKE PROTEIN RA4
8398	20936	33858	2.16	1.9E+00	P02467	SWISSPROT	COLLAGEN ALPHA Z(I) CHAIN PRECURSOR
8396	20936	33859	2.16	1.9E+00	P02467	SWISSPROT	COLLAGEN ALPHA 2(1) CHAIN PRECURSOR
8583	21132		2.45		BF360206.1	EST_HUMAN	CM3-MT0114-010900-323-h12 MT0114 Homo sapiens cDNA
8825	21364		1.35	1.9E+00	051781	SWISSPROT	ARGININE DEIMINASE (ADI) (ARGININE DIHYDROLASE) (AD)
							ab94e04.s1 Stratagene lung (#937210) Homo sepiens cDNA clone IMAGE:8545743' similar to contains Alu
9548	22048	32009	9.0	1.9E+00	AA669125.1	EST_HUMAN	repetitive element contains element L1 L1 repolitive element;
10456		35959	0.52	1.9E+00	AF248269.1	NT	Homo sapiens gag-pro-pol precursor protein gene, partial cds
3128	L			1.8E+00	P21004	SWISSPROT	PROTEIN B8 PRECURSOR
	L						Synechococcus sp. PCC7942 copper transporting P-ATPase (ctaA) and ATP synthase epsilon subunit
3154	15768	28234	2.42	1.8E+00	U04356.1	NT	(atpE) genes, complete cds
				į			Synechococcus sp. PCC7942 copper transporting P-ATPase (ctaA) and ATP synthase epsilon subunit
3154	15768	28235	2.42	1.8E+00	U04356.1	NT	(atpE) genes, complete cds
6027	18646		2.02	1.8E+00	P18502	SWISSPROT	HEDGEHOG RECEPTOR (PATCHED PROTEIN)
6253	18862	31634	2.02	1.8E+00	BF311999.1	EST_HUMAN	601897854F1 NIH_MGC_19 Hamo sapiens cDNA clone IMAGE:4127384 5'
6532	19132		1.53	1.8E+00	BF683327.1	EST_HUMAN	602139470F1 NIH_MGC_46 Homo sapiens cDNA clone IMAGE:4298272 5
6838	19428	32244	1.35	1.8E+00	BF305652.1	EST_HUMAN	601893489F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:4139038 5'
7119	19459	32274	1.08	1.8E+00	P21249	SWISSPROT	MAJOR ANTIGEN
							RETROVIRUS-RELATED POL POLYPROTEIN (CONTAINS: REVERSE TRANSCRIPTASE;
8060	20602	33512	0.81	1.8E+00	P11369	SWISSPROT	ENDONUCLEASE)
-					07470	TOBOSSINIS	RETROVIRUS-RELATED POL POLYPROTEIN (CONTAINS: REVERSE TRANSCRIPTASE;
278	L	34252				SWISSPROT	EMBRYONAL FYN-ASSOCIATED SUBSTRATE (HEFS)
9102	1				R31042.1	EST HUMAN	yh72c08.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:135278 5'
9186	1.			L	AW 880004.1	EST HUMAN	QV0-070030-070300-148-e03 OT0030 Homo sapiens cDNA
9763		L				SWISSPROT	CHITINASE D PRECURSOR
10183		L	3.78			NT.	Homo sapiens PR00530 mRNA, complete cds
10447	L		0.85			SWISSPROT	CYTIDINE DEAMINASE (CYTIDINE AMINOHYDROLASE) (CDA)
12075	24015		6.85	1.8F+00	AF314254 1	LN	Chlamydomonas reinhardtii alternative oxidase 1 (AOX1) gene, nuclear gene encoding mitochondrial protein
12163	1		4.96	L	_	INT	Rattus norvegicus Actin-related protein complex 1b (Arpc1b), mRNA
12476	1 1	30790	1.38	1.8E+00	BF212412.1	EST_HUMAN	601813714F1 NIH_MGC_54 Hamo sapiens cDNA clone IMAGE:4048251 5*

Page 16 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor		Horno sapiens chromosome 21 segment HS21C080		LEVANSUCRASE (BETA-D-FRUCTOFURANOSYL TRANSFERASE) (SUCROSE 6-FRUCTOSYL TRANSFERASE)	1	J CM0-BT0282-171299-127-e05 BT0282 Homo sapiens cDNA	Г			ORPHAN NUCLEAR RECEPTOR NR1D1 (V-ERBA RELATED PROTEIN EAR-1) (REV-ERBA-ALPHA)	Mus musculus T cell receptor gamma locus, TCR gamma 2 and gamma 4 gene clusters	Mus musculus T-cell acute lymphocytic leukemia 1 (Tal1), mRNA		Hippoglossus hippoglossus interferon inducible Mx protein (Mx) mRNA, complete cds	Г	M.musculus Ank-1 mRNA for enythroid ankyrin	M.musculus Ank-1 mRNA for enythroid ankyrin			Homo sapiens HSPC262 mRNA, partial cds	 67B7 Human retina cDNA Tsp509I-cleaved subtibrary Homo sapiens cDNA not directional 	hu82d07.x1 NCI_CGAP_Gas4 Homo saplens cDNA clone IMAGE:2257549 3' similar to contains MSR1.t1 MSR1 receitive element:	1		Homo sapiens lens epithelium-derived growth factor gene, alternatively spliced, complete cds	Homo sapiens small proline-rich protein (SPRR3) gene, exons 1, 2, and 3 and complete cds	Mus musculus ST6CatNAcIII gene, exon 2	B.napus gene encoding endo-polygalacturonase	zd25f01.r1 Soeres_feta_heart_NbHH19W Homo sapiens cDNA clone IMAGE:341689 5' similar to gb:D29805 N-ACETYLLACTOSAMINE SYNTHASE (HUMAN);	Γ
Top Hit Detabase Source	SWISSPROT	N	EST_HUMAN	TOGGSSIMS	EST HUMAN	EST_HUMAN	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	NT	NT	EST_HUMAN	LΖ	EST_HUMAN	TN	NT	SWISSPROT	SWISSPROT	NT	EST_HUMAN	FST HUMAN		EST_HUMAN	N L	N T	Z	N	EST_HUMAN	TOT TOTAL
Top Hit Acession No.	Q60114	AL163280.2	Al141067.1	0.80114	BE083546.1	BE063546.1	Q9TTRB	Q03703	Q03703	P20393	AF021335.1	6755715 NT	BF530630.1	AF245513.1	BF308000.1	X69063.1	X69063.1	060479	060479	AF161380.1	W22424.1	AI878443 1		AI198573.1	AF199339.1	AF077374.1	Y11344.1	X98373.1	W58426.1	05670077 4
Most Similar (Top) Hit BLAST E Value	1.7E+00	1.7E+00	1.7E+00	1 75+000			1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00	1.7E+00		1.7E+00	1.7E+00			1.7E+00	1.6E+00	1.6E+00	1.6E+00	1.6E+00	1.6E+00	1 85+00
Expression Signal	2.08	2.37	1.29	0.74	1.65	1.65	3.35	1.33	1.33	1.63	96.0	1.34	0.57	0.61	2.08	0.49	0.49	2.25	2.25	1.65	2.16	1.52		1.79	21.82	4.3	1.04	1.13	1.5	7.22
ORF SEQ ID NO:	26259	27458	27554	20580	31137	31138	31545	32654	32865	32693	33247	33425	33452	33933		34096	34097	34545	34548		36985	30003		30873	27228	27238	27243		28084	
Exon SEQ ID NO:	13750	14883	14979	17141	18422	18422	18780	19798	19798	19834	20339	20518	20548	21018	21101	21177	21177	24792	24792	22024	23917	24320		24659	14658	14668	14673	14894	15604	ł
Probe SEQ ID NO:	1147	2311	2411	ARRO	5797	5797	6168	7270	7270	7306	7798	7978	8008	8479	8562	8638	8638	9076	9076	8524	11487	12030		12558	2078	2087	2083	2323	2888	4104

WO 01/57277 PCT/US01/00669

Page 17 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

<u> </u>		_	_	_	-		_	_	_	_	_	_	_	_	_	_	_	_	_			_	_	_			_	_		_	
	Top Hit Descriptor	Homo sapiens proliferation-associated SNF2-like protein (SMARCA6) mRNA, complete cds	Homo sapiens proliferation-associated SNF2-like protein (SMARCA6) mRNA, complete cds	Urdeuthis chinensis cytochrome c oxidase subunit (COI) gene, mitochondrial gane encoding mitochondrial	protein, partial cds	Uroteuthis chinensis cytochrome c oxidase subunit I (COI) gene, mitochondrial gene encoding mitochondrial protein partial cds.	Mus musculus ST8GaNAcIII gene, exon 2	Mus musculus ST6GalNAcili gene, exon 2	Brachydanio rerio MHC class II DA-beta-2*01 gene, 3' end	Homo sapiens transglutaminase type I (Tgasel) gene, promoter region	IL2-UT0073-080900-145-E02 UT0073 Homo sapiens cDNA	UI-H-Bi2-ahr-b-04-0-UI:s1 NCI_CGAP_Sub4 Homo sapiens cDNA clone IMAGE:2727511 3	RC0-CT0415-200700-032-c10 CT0415 Homo sapiens cDNA	VIRULENCE FACTOR MVIN HOMOLOG	Mus musculus SIL, MAP_17, CYP_a, SCL & CYP_b genes	Homo sapiens hypothetical protein PRO0971 (PRO0971), mRNA	Homo sapiens hypothetical protein PRO0971 (PRO0971), mRNA	M.musculus COL3A1 gene for collagen alpha-l	M.muscutus COL3A1 gene for collegen alphe-	ph6b6_19/1TV Outward Alu-primed hncDNA library Homo sapiens cDNA clone ph6b6_19/1TV	Drosophila metanogaster signal transducting adaptor protein (STAM), serine threonine kinase Ial (IAL), and Ialne transfer CDN21) denes, complete cds	QV4-LT0016-090200-100-407 LT0018 Homo sepiens cDNA	QV4-LT0016-090200-100-d07 LT0016 Homo sapiens cDNA	Mus musculus T cell receptor gamma locus, TCR gamma 1 and gamma 3 gene clusters	CAPSID PROTEIN P40 [CONTAINS: ASSEMBLIN (PROTEASE); CAPSID ASSEMBLY PROTEIN]	CAPSID PROTEIN P40 [CONTAINS: ASSEMBLIN (PROTEASE); CAPSID ASSEMBLY PROTEIN]	Homo sapiens transglutaminase type I (Tgasel) gene, promoter region	Homo sapiens unknown mRNA	Rattus norvegicus jun dimerization protein 2 (jdp-2) mRNA, complete cds	Chlamydophila pneumoniae AR39, section 32 of 94 of the complete genome	Mus musculus a disintegrin and metalloproteinase domain (ADAM) 15 (metargidin) (Adam15), mRNA
	Top Hit Detabase Source	NT	Ę		L	Ę	Ż	\ V	۲	N	EST_HUMAN	EST_HUMAN	EST_HUMAN	SWISSPROT	Z	LN LN	Z	N	ΙΝ	EST_HUMAN	Į.	EST HUMAN	EST_HUMAN	¥	SWISSPROT	SWISSPROT	N	NT	Ž	NT	ΓŅ
	Top Hit Acession No.	0 AF155827.1	1.6E+00 AF155827.1		1.6E+00 AF075394.1	1 8F+00 AF075394 1	r/11344.1	Y11344.1	-04808.1	1.6E+00 AF005631.1	1.6E+00 BF380703.1	1.6E+00 AW 294881.1	1.6E+00 BE697267.1	246378	1.6E+00 AJ297131.1	11437222 NT	11437222 NT	X52046.1	X52046.1	30 T41290.1	4 8E+00 AE121381 1	1.8E+00 AW835844.1	1.8E+00 AW835644.1	1.6E+00 AF037352.1	P54817	P54817	1.6E+00 AF005631.1	1.6E+00 AF104313.1	1.5E+00 U53449.1	00 AE002201.2	6752961 NT
	Most Similar (Top) Hit BLAST E Value	1.6E+00	1.6E+00		1.6E+00	1.6F+00	1.6E+00 Y11344.1	1.6E+00 Y11344.1	1.6E+00 L04808.1	1.6E+00	1.6E+00	1.6E+00	1.6E+00	1.6E+00 Q46378	1.6E+00	1.6E+00	1.6E+00	1.6E+00 X52046.1	1.6E+00 X52046.1	1.6E+00	4 85+00	1.8E+00	1.8E+00	1.6E+00	1.6E+00 P54817	1.6E+00 P54817	1.6E+00	1.6E+00	1.5E+00	1.5E+00	1.5E+00
	Expression Signal	1.11	1.11		9.6	6	22	2.2	1.95	0.92	0.93	1.07	2.32	1.09	3.24	0.95	0.95	3.16	3.16	1.34	0.52	0.92	0.92	0.49	1.59	1.56	6.41	2.92	4.02	2.17	1.98
	ORF SEQ ID NO:		29471		30145	20148		30229		31434		32218	32680		33786		34295	33221	33222	35119	35544					<u>.</u>		37072	25173		
	Exan SEQ ID NO:	17030	17030	l	17715	17715	17807	17807	18613	18689	19193	19402	19821	20515	20861	21370	21370	24790	24790	22148	225.47	L	22583	L				24000	1	12912	13272
	Probe SEQ ID NO:	444	4444		5145	5145	5243	5243	5993	6072	8238	6811	7283	7973	8320	8831	8831	9381	9381	9649	40052	10088	10088	10246	10650	10686	10723	11552	35	252	648

Page 18 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	_	_	_	_	_				_	~	_	_	_			_	_		_	_	_	_		_			_	_	,	_	_
Тор Hit Descriptor	Mus musculus receptor pratein tyrosine phosphatase-rho (Ptprt) gene, exons 10 and 11 and partial cds	Potato virus A RNA complete genome, Isolate U	Mus musculus T-cell lymphoma invasion and metastasis 1 (Tiam1), mRNA	Potato virus A RNA complete genome, isolate U	Deinococcus radiodurans R1 section 82 of 229 of the complete chromosome 1	tt12f10.x1 NCI_CGAP_GC8 Homo sapiens cDNA clone IMAGE:2240587 3' similar to TR:000237 000237 HKF-1.;	#12f10.x1 NCI_CGAP_GC6 Homo sepiens cDNA clone IMAGE:2240587 3' similar to TR:000237 000237 HKF-1	yg10e02.r1 Sogres infant brain 1NIB Homo sapiens cDNA clone IMAGE:31693 5	601478745F1 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:3881555 5	HYPOTHETICAL 118.4 KD PROTEIN IN BAT2-DAL5 INTERGENIC REGION PRECURSOR	HYPOTHETICAL 118.4 KD PROTEIN IN BAT2-DAL5 INTERGENIC REGION PRECURSOR	ak26110.s1 Scares_testis_NHT Homo sapiens cDNA clone INAGE:14071153'	601509586F1 NIH_MGC_71 Hamo sapiens cDNA clone IMAGE:3911181 5'	Mouse germline IgM chain gene, mu-delta region	Homo sapiens hGPIb alpha gene for platelet glycoprotein Ib alpha, complete cds	601882662F1 NIH_MGC_57 Hamo sapiens cDNA clane IMAGE:4095135 5	yj03h01.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:147697 5'	QV3-CT0192-261099-008-d09 CT0192 Homo sepiens cDNA	RC0-TN0078-150900-034-g05 TN0078 Homo sepiens cDNA	602035771F1 NCI_CGAP_Bm64 Homo sapiens cDNA clone IMAGE:4183865 5'	ze38g06.r1 Soares retina N2b4HR Hamo sapiens cDNA clone IMAGE:361306.5	ze38g06.r1 Soares retina N2b4HR Homo sapiens cDNA clone IMAGE:361306 5'	DKFZp547P243_s1 547 (synonym: hfbr1) Homo sapiens cDNA clone DKFZp547P243 3'	Maize mitochondrial tRNA-Ser gene and tRNA-Phe pseudogene	Human mRNA for KIAA0146 gene, partial cds	Thermoplasma acidophilum complete genome; segment 3/5	Homo sapiens DKFZP586M0122 protein (DKFZP586M0122), mRNA	Homo sapiens DKFZP586M0122 protein (DKFZP586M0122), mRNA	yn57e03.r1 Soares adult brain N2b5HB55Y Homo sapiens cDNA clone IMAGE:172540 5'	Helicobacter pylori glutamine synthetase (glnA) gene, complete cds	Ovis aries prion protein gene, complete cds
Top Hit Database Source	IN	TN	LN	TN	TN	EST_HUMAN	FST HIMAN	EST HUMAN	EST HUMAN	SWISSPROT	SWISSPROT	EST_HUMAN	EST_HUMAN	TN	NT	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	N.	L	NT	ΙZ	N	EST_HUMAN	NT	TN
Top Hit Acession No.	AF275265.1	AJ131402.1	6678350	AJ131402.1	AE001945.1	A1655301.1	A1655301 1	R17879.1	BE785356.1	P47179	247179	AA889259.1	BE887446.1	K02138.1	AB038516.1	1.1		1.5E+00 AW375697.1	1.5E+00 BF376754.1	1.5E+00 BF337944.1	1.5E+00 AA017689.1	1.5E+00[AA017689.1	1.5E+00 AL134197.1	X07380.1	D63480.1	1.5E+00 AL445065.1	7661685 NT	7661685 NT	H19859.1	1.4E+00 AF053357.1	1.4E+00 U67922.1
Most Similar (Top) Hit BLAST E Value		1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5F+00	1=	1,5E+00	1.5E+00	1.5E+00 P47179	1.5E+00 /	1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5E+00 R81928.1	1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5E+00	1.5E+00 X07380.1	1.5E+00 D63480.1	1.5€+00	1.4E+00	1.4€+00			
Expression Signal	2.55	2.13	1.83	1.54	2.0	1 6.0	760	2.68	1.37	20.84	20.84	1.02	0.85	1.1	0.53	0.54	0.0	1.12	5.97	1.47	2.95	2.95	4.1	9.57	1.59	4.99	1.8	1.8	1.32	96.0	7.8
ORF SEQ ID NO:	27101	27592	27690	27592	28510	31250	31251	31930		32599	32600	32774	33519	34037		34528	34862	35016	35257			98538	36785		30615		25169		26909	\ \ .	
Exan SEQ ID NO:		15021	15120	15021	16029	18525	18525		19714	19745	19745	19909	20607	21117			21913	22053	ı	22460		22593	23730		25010	24465	12711	12711	14364	14888	14942
Probe SEQ ID NO:	1960	2454	2556	3172	3421	5903	5003	6538	7182	7214	7214	7383	8065	8578	8946	9061	9404	9553	9774	9965	10098	10098	11277	11416	12022	12255	32	32	1774	2316	2372

Page 19 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

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Probe SEQ ID NO:	Exan SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
2693	15250	27820	1.45	1.4E+00	X74463.1	L×	Human papillomavirus type 7 genomic DNA
2802	15354	27922	2.79	1.4E+00	AF064564.2	F	Fugu rubripes neurofibromatosis type 1 (NF1), A-kinase anchor protein (AKAP84), BAW protein (BAW), and WSB1 protein (WSB1) genes, complete cds
2802	15354	27923	2.79	1.4E+00	AF064564.2	LZ	Fugu rubripes neurofibromatosis type 1 (NF1), A-kinase anchor protein (AKAP84), BAW protein (BAW), and WSB1 protein (WSB1) genes, complete cds
3376	ł			1.4E+00	5453733 NT	N.	Homo sepiens Mad4 homdog (MAD4) mRNA
4342	16929	28369	1.14	_	AW900455.1	EST_HUMAN	CMO-NN1005-140300-286-h06 NN1005 Homo sapiens cDNA
4342	16929	28370		1.4E+00	AW900455.1	EST_HUMAN	CM0-NN 1005-140300-286-h06 NN 1005 Homo sapiens cDNA
4685	17267		1.78	1.4E+00	BF681547.1	EST_HUMAN	602156687F1 NIH_MGC_83 Homo sapiens cDNA clone IMAGE:4297556 5'
5275	18206	30657	1.76	1.4E+00	AW054976.1	EST_HUMAN	wt45g07.x1 NCI_CGAP_Pan1 Homo saplens cDNA clone IMAGE:2510460 3'
5719			5.04	1.4E+00	AB032983.1	NT	Homo sapiens mRNA for KIAA1157 protein, partial cds
6425		31811	2.73	1.4E+00	Q13472	SWISSPROT	DNA TOPOISOMERASE III ALPHA
6437	25116		4.4	1.4E+00	AB020712.1	NT	Homo sapiens mRNA for KIAA0905 protein, complete cds
6544	19143	31936	2:32	1.4E+00	Q92777	SWISSPROT	SYNAPSIN II
6544	19143	31937	2:32	1.4E+00	<i>Q92777</i>	SWISSPROT	SYNAPSIN II
6583	19181	31981	19'0	1.4E+00	11096333 NT	N	Mus musculus WW domain binding protein 11 (Wbp11-pending), mRNA
6911	19570	32398	77.0	1.4E+00	AW893057.1	EST_HUMAN	CM3-NN0008-300300-132-b12 NN0006 Homo sapiens cDNA
7330	19857	32720	2.31	1.4E+00	AJ133269.1	TN	Homo saplens caveolin-1/-2 locus, Contig1, D7S522, genes CAV2 (exons 1, 2a, and 2b), CAV1 (exons 1 and 2).
7343	I _			1 45+00	AW487780 4	MAN II	he23t05.x1 NCI_CGAP_CML1 Homo sepiens cDNA clone IMACE:2019873 3' similar to contains Alu
				3			CHILDRAN ASE DEFOLISOD OF LICAN 4.4 ALDER CHILDRAN
8277	20818		0.68	1.4E+00	P07683	SWISSPROT	GLUCOHYDROLASE)
8728	21268		4.01		AJ271735.1	NT	Homo sapiens Xq pseudoautosomal region; segment 1/2
9023	21560		2.13	1.4E+00	R20459.1	EST_HUMAN	yg33f12.r1 Soares infant brain 1NIB Homo sapiens cDNA clone IMAGE:34345 5'
9125			3.72	1.4E+00	BE064667.1	EST_HUMAN	RC1-BT0313-301299-012-f05 BT0313 Homo sapiens cDNA
9158	21693		85.0	1.4E+00	AF134844.1	IN	Sceloporus undulatus ornithine transcarbamylase (OTC) mRNA, complete cds
10109	22604	35594	77.0	1.4E+00	BF575545.1	EST_HUMAN	602133135F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4288137 5'
10150	22645	35637	0.67	1.4E+00	BE145374.1	EST_HUMAN	IL5-HT0198-291099-008-C04 HT0198 Homo saplens cDNA
10150	22645	35638	0.67	1.4E+00	BE145374.1	EST HUMAN	IL5-HT0198-291099-008-C04 HT0198 Homo sapiens cDNA
10418	22912		1.11	1.4E+00	D63441.1	NT	Pandorina colemaniae chloroplast rbcL gene for ribulose bisphosphate carboxylase, partial cds
10418	22912	35913	1.11	1.4E+00	D63441.1	NT	Pandorina colemaniae chloropiast rocL gene for ribulose bisphosphate carboxylase, partial cds
10948	23463	36485	2.16	1.4E+00	AA195528.1	EST HUMAN	z 36609.r1 Soares_NhHMPu_S1 Homo sapiens cDNA clone IMAGE:665512 5' similar to contains element MER22 repetitive element ;
	ı	١					

Page 20 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

					,		
Probe SEQ 1D NO:	SEQ 1D NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
11104	23614	36654	6.28	1.4E+00	AB006682.1	N	Homo sapiens APECED mRNA for AIRE-1, complete cds
11283	23736	36791	3.92	1.4E+00	BE962107.2	EST_HUMAN	801655184R1 NIH_MGC_65 Homo sapiens cDNA clone IMAGE:3845805 3'
11283	23736	36792	3.92	1.4E+00	BE962107.2	EST_HUMAN	601655184R1 NIH_MGC_65 Homo sapiens cDNA clone IMACE:3845805 3'
11304	23797	36855	3.19	1.4E+00	U30790.1	L	Pneumocystis cerinii f. sp. ratti guanine nucleotide binding protein alpha subunit (pog 1) gene, complete cds
11304	23797	36856	3.19	1.4E+00	U30790.1	LN	Pneumocystis carinii f. sp. ratti guanine nuclectide binding protein alpha subunit (pcg1) gene, complete cds
11865	24935			1.4E+00	AL161500.2	NT	Arabidopsis thaliana DNA chromosome 4, contig fregment No. 12
12267	25108		2.38	1.4E+00	11545836 NT	NT	Homo sapiens cutaneous T-cell lymphorna tumor antigen se70-2 (SE70-2), mRNA
985	13225		1.38	1.3E+00	Z73640.1	NT	M.mucedo gene encoding 4-Dihydromethyl-trisporate dehydrogenase
935	13548	26065	2.33	1.3E+00	AJ271192.1	NT	Cantharellus sp. partial 25S rRNA gene, isolate Tibet
1168	13770		22.19	1.3E+00	Y19213.1	NT	Homo sapiens putative psihHbA pseudogene for hair keratin, exons 2 to 7
1340	13935	26456	13.67	1.3E+00	TN 8867988	Į,	Homo sapiens zinc finger protein 157 (HZF22) (ZNF157) mRNA
1340	13935		13.67	1.3E+00	TN 8687054	IN	Homo sapiens zinc finger protein 157 (HZF22) (ZNFJ57) mRNA
140	13994		1.05	1.3E+00	U61730.2	IN	Coix lacryma-jobi dihydrodipicolinate synthase (dapA) gene, complete cds
1653	14245		2.35	1.3E+00	AE002338.2	NT	Chlamydia muridarum, section 66 of 85 of the complete genome
							Cyprinus carpio MRPb and MASPb genes for mannose-binding lectin-associated serine protease (MASP)
2285	14859		1.1		AB030447.1	NT	and MASP-related protein, complete cds
2586	15149		76.0	1.3E+00	BE966735.2	EST_HUMAN	601681233R1 NIH_MGC_72 Home sapiens cDNA clone IMAGE:3915945 3'
2968	15581	28060	99'0	1.3E+00	6755821 NT	N	Mus musculus alpha-spectrin 1, erythroid (Spna1), mRNA
							Fugu rubripes gamma-aminobutyric acid receptor beta subunit gane, partial cds; 55kd erythrocyte membrane
							protein (P55), synaptic vesicle-associated integral membrane protein (VAMP-1), procollagen C-proteinase
3657	16260	28732	0.91	1.3E+00	AF016494.1	۲	enhancer protein (PCOLCE) genes, complete c>
4713		L		1.3E+00	6755621 NT	NT	Mus musculus alpha-spectrin 1, erythrold (Spna1), mRNA
5184	L_		0.92	1.3E+00	AJ252087.1	N	Plasmodium reichenowi partial 83/AMA-1 gene for apical membrane antigen 1
5184			0.92	1.3E+00	AJ252087.1	۲	Piasmodium reichenowi partial 83/AMA-1 gene for apical membrana antigen 1
5705	Ι.		1.06		P19732	SWISSPROT	PHENOL HYDROXYLASE P3 PROTEIN (PHENOL 2-MONOOXYGENASE P3 COMPONENT)
6169		L	7.47	L	AW362834.1	EST HUMAN	PMC-CT0289-291189-004-f08 CT0289 Homo sapiens cDNA
6169	L	31547	7.47			EST_HUMAN	PM0-CT0289-291199-004-f08 CT0289 Homo sapiens cDNA
8549	19147		1.24	1.3E+00	M33496.1	Ā	D.melanogaster no-on-transient A gene product, complete cds
6847	19437		0.71	L	Q00156	SWISSPROT	HYPOTHETICAL GENE 64 PROTEIN
6926	19585	32415	0.85	1.3E+00	M13918.2	NT	Homo sapiens fibronectin receptor alphe-subunit precursor (ITGA5) mRNA, partial cds
7033	19567	32394	1.01	1.3E+00	BE538819.1	EST_HUMAN	601061420F1 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3447965 5'

Page 21 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Database Source	EST_HUMAN	40 SWISSPROT PHOSPHOHYDROLASE)	N.	EST_HUMAN	EST_HUMAN		EST_HUMAN	EST_HUMAN	EST_HUMAN	2084.1 NT Homo sapiens heparan glucosaminy N-deacetylasaN-sulfotransferase-2 gene, complete cds	N	Z	LN⊤	LYSOSOMAL ALPHA-MANNOSIDASE PRECURSOR (MANNOSIDASE, ALPHA B) (LYSOSOMAL ACID	EST LIMAN	NIMONIC TO THE	2 5	IN LOS	33379.2 EST_HUMAN 601657145KT NIH_MIGC_07 Home septens CURN clone image; 3000183.3	, L	NT NT	23637 NT	EST_HUMAN	1 EST_HUMAN	SWISSPROT	S99 SWISSPROT MRNA 3'-END PROCESSING PROTEIN RNA15	INT	EST_HUMAN	NT	NT	37873.1 NT Cavia porcellus inwardly-rectifying potassium channel Kir2.2 (KCNJ12) gene, complete cds
Top Hit Acession No.	3E+00 BE243571.1		12.1	1.3E+00 BE963379.2	1.3E+00 BE974280.1	9910247	1.3E+00 AI927629.1		1.3E+00 H42881.1	1.3E+00 AF042084.1	1.3E+00 X72019.1			000 or					1.3E+00 BE963379.2		2	3837	1.3E+00 H42881.1	1			П	11.1			3.1
Most Similar (Top) Hit BLAST E Value	1.3E+00	1.3E+00 P24540	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00 H42881.1	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1 25 +00	1.35+00	1.35400	1.35+00	1.3E+00	1.35 +00	13F+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00 Q14117	1.3E+00 P25299	1.3E+00	1.3E+00	1.3E+00	1.3E+00	1.3E+00
Expression	0.79	3.97	2.08	2.54	0.89	1.57	98.0	0.48	0.48	4.54	2.12	2.12	1.1	. 63	70.	17.	0.03	0.83	3.85	2 41	0.65	0.52	0.48	0.48	4.66	2.3	2.17	1.87	3.09	3.09	3.63
ORF SEQ ID NO:	32633	32868			33856		34184		34541		34917			0,000					35263				36008	36009		36316			36926		
Exan SEQ ID NO:	19889	20003	20780	<u>i_</u>	L	21183	21264			21960	Ι.	I	L.	l	20100	1		\perp		22825	1	1	l	l	1		1_	L		Ш	24312
Probe SEQ ID NO:	7157	7481	8238	8384	8496	8644	8725	9073	9073	9434	9443	9443	9542	2	000	200	9740	9740	9780	10114	10478	10504	10507	10507	10573	10785	10807	11215	11414	11488	12011

Page 22 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	802023185F1 NCI_CGAP_Bm67 Home sapiens cDNA clone IMAGE:4158452 5'	E1 GLYCOPROTEIN PRECURSOR (MATRIX GLYCOPROTEIN) (MEMBRANE GLYCOPROTEIN)	Sturnira lilium cytochrome b gene, complete cds; mitochondrial gene for mitochondrial product	Homo sapiens chromosome 21 segment HS21 C083	zi22d08.s1 Soares fetal liver_spleen_INFLS_S1 Homo sapiens cDNA clone IMAGE:431535.3'	HISTIDINE-RICH PROTEIN PRECURSOR (CLONE PFHRP-III)	HISTIDINE-RICH PROTEIN PRECURSOR (CLONE PFHRP-III)	HISTIDINE-RICH PROTEIN PRECURSOR (CLONE PFHRP-III)	Homo sepiens hypothetical protein PRO3077 (PRO3077), mRNA	Elaeis oleifera sesquiterpene synthase mRNA, complete cds	pea seed-borne mosaic virus complete genome	pea seed-barne masaic virus complete gename	Homo sapiens G-protein coupled receptor 14 (GPR14) gene, complete cds	Homo saplens post-synaptic density 95 (DLG4) gene, complete cds	Homo sapiens mRNA for KIAA0874 protein, partial cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 63	Arabidopsis thaliana DNA chromosome 4, contig fregment No. 63	CONJUGAL TRANSFER PROTEIN TRBE PRECURSOR	Homo sapiens LHX3 gene, intron 2	Mus musculus subtilisin-like serine protease LPC (PC7) gene, exons 1 to 9, partial cds	MRo-FT0175-050900-203-g06_1 FT0175 Hamo sapiens cDNA	Homo sapiens LHX3 gene, Intron 2	Rattus rattus cardiac AE3 gene, exons 1-23	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 21	Homo sapiens post-synaptic density 95 (DLG4) gene, complete cds	T.pinnatum chloroplast rbcl. gene, partial	G.gailus T-cadherin mRNA, complete cds	Human extracellular calcium-sensing receptor mRNA, complete cds	MR3-ST0191-140200-013-c05 ST0191 Homo sapiens cDNA	Homo sapiens zinc finger protein ZNF191 (ZNF191) gene, complete cds	D.hydei ayr repeat cluster DNA, fragment D	QV4-BN0090-270400-190-e03 BN0090 Homo sepiens cDNA	C.glutamicum pta gene and ackA gene	Cightamtcum pta gene and eckA gene
Top Hit Database Source	EST_HUMAN	SWISSPROT	Z	Į.	EST_HUMAN	SWISSPROT	SWISSPROT	SWISSPROT	LN	TN	LN	LΝ	TN	TN	LN	LN	ΤN	SWISSPROT	ΙNΤ	IN	EST_HUMAN	IN	LN.	NT	<u>ا</u> ع	LN TA	Z	F	EST_HUMAN	NT	LN L	EST_HUMAN	NT	NT
Top Hit Acession	BF348043.1	P33464	AF187035.1	AL163283.2	AA676246.1	P05228	P05228	P05228	8924234	AF080245.2	AJ252242.1	AJ252242.1	AF140631.1	AF156495.1	AB020681.1	AL161563.2	AL181563.2	P54910	AF188740.1	U75902.1	BF373570.1	AF188740.1	M87050.1	AL161509.2	AF156495.1	Y09200.1	M81779.1	U20760.1	AW813276.1	AF016052.1	X74885.1	BE003113.1	X89084.1	X89084.1
Most Similar (Top) Hit BLAST E Value	1.3E+00		1.3E+00	1.3E+00	1,2E+00	1.2E+00	1.2E+00			1.2€+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2€+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2€+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00	1.2E+00		1.2E+00	1.2E+00	1.2E+00	1.2E+00
Expression Signal	3.47	1.76	2.08	1.25	9.75	2.5	1.04	1.04	1.9	4.87	1.3	1.3	53.59	1.53	1.16	71.17	71.7	3.43	0.57	89.8	1.87	1.12	2.09	1.08	1.5	9.41	0.77	1.08	2.27	0.72	2.17	3.98	1.43	1.43
ORF SEQ ID NO:	30949				25784	25983	25984	25985		26316	26361	26362	27208	27559			28288		28489	28839		28489		29623	L			30729	31077	31395	31879	31743		31823
Exon SEQ ID NO:	24423	24828	24500	24904	13302	13472	13472	13472	13524	13803	13844	13844	14835	14985	15758	15813	15813	15935	16007	16374	16655	16007	17153	17204	17241			18258	18369		18908	18965		Ш
Probe SEQ ID NO:	12192	12204	12303	12673	878	858	858	858	911	1203	1247	1247	2054	2417	3144	3201	3201	3325	3399	3774	4058	4386	4570	4621	4659	4690	4791	5629	5743	6034	6300	6361	6433	6433

WO 01/57277 PCT/US01/00669

Page 23 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID 8	Exan SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acessian No.	Top Hit Datebase Source	. Top Hit Descriptor
6475	19076	31859	34.96	1.2E+00	AA759254.1	EST_HUMAN	ah84g12.s1 Soares_testis_NHT Homo saplens cDNA clone 1322374 3'
6829	19225	32030	2.25	1.2E+00	AW813276.1	EST_HUMAN	MR3-ST0191-140200-013-c05 ST0191 Homo saplens cDNA
6995	19493	32314	1.18	1.2€+00	AB029010.1		Homo sapiens mRNA for KIAA1087 protein, partial cds
7007	19505		2.8	1.2E+00	AJ002141.1	NT	Mus musculus DSPP gene
7300	19828		9.0	1.2E+00	AJ271735.1	NT.	Homo saplens Xq pseudoautosomal region; segment 1/2
7417	24782	32808	1.59	1.2E+00	AV734585.1	T_HUMAN	AV734585 cdA Homo saplens cDNA clone cdAAFH03 5'
7848	20158	33045	2.84	1.2E+00	X74207.1	LN	Lilectis pyrD and pyrF genes
8504	21043		3.05	1.2E+00	AB033030.1	١	Homo sapiens mRNA for KIAA1204 protein, pertial cds
							ALPHA, ALPHA-TREHALOSE-PHOSPHATE SYNTHASE [UDP-FORMING] 123 KD SUBUNIT
8597	21136	34051	0.69	1.2E+00	P38427	SWISSPROT	(TREHALOSE-APHOSPHATE SYNTHASE) (UDY-GLUCUSE-GLUCUSEPHUSPHATE GLUCOSYLTRANSFERASE)
6088	21348		0.53	1.2E+00	7706271	LN.	Homo sepiens CGI-30 protein (LOC51611), mRNA
8955	21493	34418	2.03	1.2E+00	AW377210.1	EST_HUMAN	MR2-CT0222-201089-001-607 CT0222 Homo saplens cDNA
9319	21833		2.92	1.2E+00	232850.1	L	R.communis gene for pyrophosphate-dependent phosphofructokinase beta subunit
9523	22023	34981	1.86	1.2€+00	D11745.1	EST_HUMAN	HUMHM01A01 Liver HepG2 cell line. Homo sapiens cDNA clone hm01a01
9844	22342	35324	3.47	1.2E+00	X56832.1	NT	H.saplens ENO3 gene for muscle specific endase
10229	22724		29'0	1.2E+00	AB009666.1	NT	Homo sapiens klotho gene, exon 1
11224	23755	36813	2.19	1.2€+00	AW817817.1	EST_HUMAN	PM0-ST0264-161199-001-d01 ST0264 Home saplens cDNA
11282	23790		6.64	1.2E+00	BE160781.1	EST_HUMAN	PM1-HT0422-160200-007-g10 HT0422 Homo sapiens cDNA
11331	23029	_	3.78	1.2E+00	US0147.1	NT	Rattus norvegicus synapse-associated protein 102 mRNA, complete cds
11976	24907		32.4	1.2E+00	AL163203.2	INT	Homo sapiens chromosome 21 segment HS21C003
11998	24304		2.11	1.2E+00	AP001515.1	NT	Bacillus halodurans genomic DNA, section 9/14
489	13122	25608	1.19	1.1E+00	D86980.1	NT	Human mRNA for KIAA0227 gene, partial cds
1799	14389	26934	1.48	1.1E+00	AW995393.1	EST_HUMAN	QV0-BN0042-170300-163-g12 BN0042 Homo saplens cDNA
2617	15179	27748	1.09	1.15+00	AF067124.1	NT	Wheat yellow mosaic virus RNA1 270 kDa precursor protein gene, complete cds
3373	15981	28458	8.32	1.1E+00	AL163213.2	NT	Homo saplens chromosome 21 segment HS21C013
3373	15981	28459	9.32	1.1E+00	AL163213.2	INT	Homo sapiens chromosome 21 segment HS21C013
3533	16138	28620	0.08	1.1E+00	8922641 NT	·	Homo sapiens hypothetical protein FLJ10749 (FLJ10749), mRNA
3830	18242	28718	1 08	1 15+00	A1808360 1	NAMIH TSE	wf54h11.x1 Soares_NFL_T_GBC_S1 Homo saplens cDNA clone IMAGE:2359481 3' similar to SW:PS31_HUMAN Q12888 PS3-BINDING PROTEIN 53BP1 ;
3781	16381			1	AE003888.1		Xyella fastidiosa, section 32 of 229 of the complete genome
3781	16381				AE003888.1	NT	Xyiella fastidiosa, section 32 of 229 of the complete genome
3889	16488			1.1E+00	XB5374.1	NT	H.parahaemolyticus hphlM(A), hphlM(C), hphlR and menB geneo
4016	16614	29087	0.67	1.1E+00	8922641 NT	Ä	Homo sapiens hypothetical protein FLJ10749 (FLJ10749), mRNA

Page 24 of 526 . Table 4 Single Exon Probes Expressed in Fetal Liver

WO 01/57277

Page 25 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

		_	_		_	_	_	_	_			_		_	_	- 7	_	_	_	_	_	- 1						_,	
	Top Hit Descriptor	Homo sapiens mRNA for KIAA0934 protein, partial cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 27	Mus musculus guanine nucleotide binding protein (G protein), gamma 3 subunit (Gng3), mRNA	DNA MISMATCH REPAIR PROTEIN MUTS	Homo sapiens KIAA0626 gene product (KIAA0628), mRNA	Klebsormidium fluitans cytochrome c oxidase subunit 2 (cox2) gene, mitochondrial gena encoding	mitochondrial protein, partial cds	Homo sapiens hypothetical protein FLJ11280 (FLJ11280), mRNA	Petroselinum crispum cytosolic glucose-6-phosphate dehydrogenase 1 (cG6PDH1) mRNA, complete cds	Petroselinum crispum cytosolic glucose-4-phosphate dehydrogenase 1 (cG6PDH1) mRNA, complete cds	wf76e11.x1 Sogres_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2381548 3'	LOW TEMPERATURE ESSENTIAL PROTEIN	Taenia solium Immunogenic protein Ts76 mRNA, partial cds	Dictyostelium discoldeum isopentenyl pyrophosphate (somerase (Dipi) mRNA, complete cds	Xenopus laevis rhodopsin gene, complete cds	Cavia cobaya mRNA for serine/threoine kinase, complete cds	Marchantia polymorpha genes for 26S rRNA, 5S rRNA, 18S rRNA, 5.8S rRNA and 26S rRNA	Girardia lignina mRNA for homeodomain transcription factor (so gene)	Hamo sepiens chromosome 21 segment HS21C018	Aedes aegypti mucin-like protein MUC1 mRNA, complete cds	V.carteri Algal-CAM mRNA	Plautia stall intestine virus RNA for nonstructural polyprotein, capsid protein precursor, complete cds	DNA GYRASE SUBUNIT B	DNA GYRASE SUBUNIT B	3-OXO-5-ALPHA-STEROID 4-DEHYDROGENASE 1 (STEROID 5-ALPHA-REDUCTASE 1) (SR TYPE 1)	3-OXO-5-ALPHA-STEROID 4-DEHYDROGENASE 1 (STEROID 5-ALPHA-REDUCTASE 1) (SR TYPE 1)	HYPOTHETICAL 67,9 KD PROTEIN C6F12.08C IN CHROMOSOME I	ef28g08.s1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone IMAGE:1032830 3' similar to WP:C42D8.3 CE04204 ;contains element MER22 MER22 repetitive element ;
Top Hit	Database Source	Ę	F	LΝ	SWISSPROT	IN		L L	٥	Ţ	LΝ	EST_HUMAN	SWISSPROT	NT	ΙN	NT	TN	IN	NT.	NT	NT	NT	ź	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	SWISSPROT	EST_HUMAN
Ton Hit American	Ö	1.1E+00 AB023151.1	1.1E+00 AL181515.2	6754021 NT	273769	11067364 NT		1.1E+00 AF068942.1	R922973 NT	1.1E+00 AF012862.1	1.1E+00 AF012862.1	1.1E+00 AI809699.1	907866	1.1E+00 AF216696.1	1.1E+00 AF234169.1	J23808.1	D88425.1	1.0E+00 AB021684.1	1.0E+00 AJ251860.1	1.0E+00 AL163218.2	1.0E+00 AF125984.1	X80416.1	1.0E+00 AB006531.1	30 P48355	P48355	P24008	00 P24008	00 014228	1.0E+00 AA628453.1
Most Similar		1.1E+00/	1.1E+00/	1.1E+00	1.1E+00 P73769	1.1E+00		1.1E+00	1.1E+00	1.1E+00	1.1E+00	1.1E+00	1.1E+00 P07866	1.1E+00	1.1E+00	1.0E+00 U23808.1	1.0E+00 D88425.1	1.0E+00 /	1.0E+00	1.0E+00	1.0E+00	1.0E+00 X80416.1	1.0E+00		1.0E+00 P48355	1.0E+00 P24008	1.0E+00	1.0E+00	1.0E+00
Fyrestin	Signal	1.59	4.82	19.39	-	2.83		4.08	5.28	3.76	3.76	6.02	1.82	2.25	1.64	3.22	3.48	2.14	1.53	7.12	0.89	1.73	0.91	1.2	1.2	4.47	4.47	0.83	0.91
0.00	Ö NÖ:	35226		35381						36526	36527	36822		30997			25271		L	25813			26928				27987	١.	28324
Exon	SEO IO	22245	22348	22406	22889	23067		23121	18028	23497	23497	23765	24275	24335	24903	12779	12789	13076	13231	13326	13328	15441	14384		<u> </u>		15517	l	l l
Probe	SEO ID	9747	9850	6066	10395	10530		10586	10978	10983	10983	11234	11948	12051	12184	103	118	443	602	705	707	1429	1794	2526	2526	2800	2900	2994	3232

Page 26 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Xenopus laevis rhodopsin gene, complete cds	Agartcus bisparus mRNA for tyrosinase	Homo sapiens calcium channel alpha (E subunit (CACNA1E) gene, exons 7-49, and partial cds, atternatively soliced	Homo saciens hypothetical protein FI J10139 (FI J10139) mRNA	Rattus norvegicus mRNA for N-acetylglucosaminytransferase III, complete cds	Pilot whale morbilivirus phosphoprotein (P) gene, partial cds	Oncorhynchus mykiss stit mRNA for rhamnose binding lectin STL1, complete cds	Hordeum vulgare gene encoding cysteine proteinase	Bos taurus micromolar calcium activated neutral protesse 1 (CAPN1) gene, exons 11-20, and partial cds	Bos taurus micromolar calcium activated neutral protesse 1 (CAPN1) gene, exons 11-20, and partial cds	Arabidopsis thaliana DNA chromosome 4, ESSA I FCA contig fragment No. 8	FIBER PROTEIN	UI-H-BI3-aix-d-09-0-UI.s1 NCI_CGAP_Sub5 Hamo sapiens cDNA clane IMAGE:3068969 3'	Mus musculus subtilisin-like serine protease LPC (PC7) gene, exons 1 to 9, partial cds	Homo sapiens cell cycle protein (PA2G4) gene, exons 2 though 5	SRB-11 PROTEIN	V.carteri gene encoding volvoxopsin	insulin-like growth factor-binding protein 4 [cattle, pulmonary artery endothellal cells, mRNA, 2028 nt]	B-CELL RECEPTOR CD22 PRECURSOR (LEU-14) (B-LYMPHOCYTE CELL ADHESION MOLECULE)	(BL-CAM)	Homo sapiens endothelin-converting enzyme 2 (ECE2) mRNA, complete cds	ac79b08.s1 Stratagene lung (#937210) Homo sapiens cDNA clone IMAGE:868791.3'	601443950F1 NIH_MGC_65 Hamo sapiens cDNA clone IMAGE:3848005 5'	601443950F1 NIH_MGC_65 Homo sapiens cDNA done IMAGE:3848005 5'	Rattus norvegicus mRNA for N-acety/glucosaminy/transferase III, complete cds	PEROXISOMAL HYDRATASE-DEHYDROGENASE-EPIMERASE (HDE) (MULTIFUNCTIONAL BETA-OXIDATION PROTEIN) (MFP) [INCLUDES: 2-ENOYL-COA HYDRATASE; D-3-HYDROXYACYL COA DEHYDROGENASE]
Top Hit Database Source	NŢ	NT NT	FIX	FX	Į.	Į.	NT	NT	LN	LΝ	NT	SWISSPROT	EST_HUMAN	F	IN	SWISSPROT	NT	L		SWISSPROT	NT	EST_HUMAN	EST_HUMAN	EST_HUMAN	ΙZ	SWISSPROT
Top Hit Acession No.	U23808.1	DAJ223816.1	AE222301 1	R022245 NT	210852.1	1.0E+00 AF200817.1	AB039022.1	297022.1	1.0E+00 AF248054.1	1.0E+00 AF248054.1	0 Z97341.2	204501	1.0E+00 AW452782.1	U75902.1	1.0E+00 AF104669.1	P46506	Y11204.1	1.0E+00 S52770.1		P20273	1.0E+00 AF192531.1	1.0E+00 AA775191.1	1.0E+00 BE868267.1	1.0E+00 BE868267.1	1.0E+00 D10852.1	Q02207
	1.0E+00	1.0E+00[/	1004301		1.0E+00 D10852.1	1.0E+00/	1.0E+00	1.0E+00 Z97022.1	1.0E+00	1.0E+00	1.0E+00	1.0E+00 P04501	1.0E+00	1.0E+00 U75902.1	1.0E+00	1.0E+00 P46506	1.0E+00 Y11204.1	1.0E+00		1.0E+00 P20273	1.0E+00	1.0E+00	1.0E+00	1.0E+00	1.0E+00	1.0E+00 Q02207
Expression Signal	0.78	1.55	ų,	28.0	17.2	9.0	-	2.56	4.54	4.54	1.22	4.41	1.58	1.79	0.83	1.5	1.27	1.22		8.58	1.36	7.92	1.49	1.49	1.28	2.1
ORF SEQ ID NO:		28813	08.500				30394	30527	31368	31369	31480	31641	31645	32018	32062		32442	32573			23093	33107	33349	33350		33753
Exon SEQ ID NO:		16345	18728	18040		17900	18010	18120	18633	18633	18727	18871	18877	19212	19258	19336	19608	19724		20016	20206		20444	28 44 44 44	17734	20832
Probe SEQ ID NO:	3659	3744	77.77	4382	5165	5339	5432	5486	6013	6013	6111	6263	6269	6615	6662	6742	6874	7192		7493	7697	7710	7902	7902	8084	8291

WO 01/57277 PCT/US01/00669

Page 27 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

					,[
Probe SEQ ID NO:	SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Vatue	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
828-1	20832	33754	2.1	1.0E+00	002207	SWISSPROT	PEROXISOMAL HYDRATASE-DEHYDROGENASE-EPIMERASE (HDE) (MULTIFUNCTIONAL BETA- OXIDATION PROTEIN) (MFP) [INCLUDES: 2-ENOYL-COA HYDRATASE ; D-3-HYDROXYACYL COA DEHYDROGENASE]
8413			0.85	1.0E+00	P51784	SWISSPROT	UBIQUITIN CARBOXYL-TERMINAL HYDROLASE 11 (UBIQUITIN THIOLESTERASE 11) (UBIQUITIN- SPECIFIC PROCESSING PROTEASE 11) (DEUBIQUITINATING ENZYME 11)
8447	20987	33902	0.48	1.0E+00	Q9Y5T5	SWISSPROT	UBIQUITIN CARBOXYL-TERMINAL HYDROLASE 16 (UBIQUITIN THIOLESTERASE 16) (UBIQUITIN- SPECIFIC PROCESSING PROTEASE 16) (DEUBIQUITINATING ENZYME 16) (UBIQUITIN PROCESSING PROTEASE UBP-M)
8447	20987	33903	0.48	1.0E+00	Q9Y5T5	SWISSPROT	UBIQUITIN CARBOXYL-TERMINAL HYDROLASE 16 (UBIQUITIN THIOLESTERASE 16) (UBIQUITIN- SPECIFIC PROCESSING PROTEASE 16) (DEUBIQUITINATING ENZYME 16) (UBIQUITIN PROCESSING PROTEASE UBP-M)
8475	24791		2,17	1.0E+00	BE147331.1	EST_HUMAN	RC1+HT0229-181099-011-e06 HT0229 Homo sapiens cDNA
8513	24052	33974	106	1.05+00	U42720 2	·	Strains immunodeficiency virus Geg protein (geg) gene, complete cds; Poi protein (pol) gene, partial cds; and Vif protein (vif), Vpr protein (vpr), Tat protein (tat), Rev protein (rev), Vpu protein (vpu), Env protein (env), and Nef protein (nef) genes. >
8659	1			L	M38427.1	ΙZ	Human immunodeficiency virus type 1 (HIV-1), Isolate SF33,
9185	21712	34655		1.0E+00	BE907592.1	EST_HUMAN	601497581F1 NIH_MGC_70 Hamo sapiens cDNA clone IMAGE:3899421 5'
9402	21911	34860	1.34	1.0E+00	6753429 NT	Į.	Mus musculus chloride channel calcium activated 1 (Clca1), mRNA
9402	21911	34861	1.34	1.0E+00	8753429 NT	NT	Mus musculus chloride channel calcium activated 1 (Cloa1), mRNA
9528		34987	2.08	1.0E+00	AV689554.1	EST_HUMAN	AV889554 GKC Horno sepiens cDNA clone GKCCYA11 5'
9534			1.33	1.0E+00	U44952.1	NT	Xenopus laevis zona pellucida C glycoprotein precursor (xIZPC) mRNA, complete cds
9534	22034	34994	1.33	1.0E+00	U44952.1	LN	Xenopus laevis zona pellucida C giycoprotein precursor (xIZPC) mRNA, complete cds
9767			0.5		X15498.1	TN	Human Coronavirus gene for membrane protein
9767			9.0	1.0E+00	X15498.1	NT	Human Coronavirus gene for membrane protein
10021	22518		0.62			NT	Homo sepiens MHC binding factor, beta (MHCBFB) mRNA
10021	22516	35511	0.62	1.0E+00	5174562 NT	TN	Homo sapiens MHC binding factor, beta (MHCBFB) mRNA
10105		35592	0.75	1.0E+00	AI077920.1	EST_HUMAN	oy15d07.s1 Soares_senescent_fibroblasts_NbHSF Homo sapiens cDNA clone IMAGE:18659013'
10230		35716	4.17	1.0E+00	AV758825.1	EST_HUMAN	AV758825 BM Homo sepiens cDNA clone BMFAWC04 5'
10372	22868	35859	19.78	1.0E+00	AA004982.1	EST_HUMAN	zh94e02.r1 Soares fetal liver_spleen_1NFLS_S1 Homo sapiens cDNA clone IMAGE:428908 5
10372		35860	19.78	1.0E+00	AA004982.1	EST_HUMAN	zh94802.r1 Soares_fetal liver_spleen_1NFLS_S1 Homo squiens cDNA clone IMAGE:428906 5'
10404	22898	35893	0.93	1.0E+00	L11910.1	LN	Human retinoblastoma susceptibility gene exons 1-27, complete cds
10853	23374	38383	1.87	1.0E+00	S90825.1	TN	PBR1=proline-rich protein (intron 3) [human, Genomic, 898 nt]
11587	i	30527	1.57	1.0E+00	297022.1	LN	Hordeum vulgare gene encoding cysteine proteinase
11837	24201		4.85		1.0E+00 P15308	SWISSPROT	THROMBOMODULIN PRECURSOR (FETOMODULIN) (TM)

Page 28 of 526 Table 4

Single Exon Probes Expressed in Fetal Liver	Top Hit Descriptor	EST382293 MAGE resequences, MAGN Homo sapiens cDNA	Drosophila melanogaster regulator of G-protein signalling LOCO III mRNA, complete cds	Drosophila melanogaster regulator of G-protein signalling LOCO III mRNA, complete cds	Homo sapiens chromosome 21 segment HS21C102	Apple mosaic virus RNA 2 putative polymerase gene, complete cds	SERINE/THREONINE PROTEIN KINASE MINIBRAIN	PROBABLE OXIDOREDUCTASE ZK1290.5 IN CHROMOSOME II	Lycopersicon esculentum purative Mi1 copy 1 nematode-resistance gene	B2 BRADYKININ RECEPTOR (BK-2 RECEPTOR)	Danio rerio mRNA for Eph-like receptor tyrosine kinase rtk8	AMINO-ACID ACETYLTRANSFERASE (N-ACETYLGLUTAMATE SYNTHASE) (AGS) (NAGS)	Calithrix jacchus UBE1 gene derived retroposon on the Y chromosome	Xenopus laevis rac GTPase mRNA, complete cds	PROBABLE ENDONUCLEASE IV (ENDODEOXYRIBONUCLEASE IV)	601653583R2 NIH_MGC_55 Homo sapiens cDNA clone IMAGE:3838461 3'	601653583R2 NIH_MGC_55 Homo sapiens cDNA clone IMAGE:3838461 3'	Enterobacteriaceae sp. JM983 partial groES gene for GroES-like protein and partial groEL gene for GroEL- like system (solide, 1M693)	process, isolate divisors	Enterobacteriaceae sp. JM983 partial groES gene for GroES-like protein and partial groEL gene for GroEL. like protein, isolate JM983	601456337F1 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:3880049 5	601456337F1 NIH_MGC_68 Hamo sepiens cDNA clone IMAGE:3860049 5	IOSPHOGLUCOMUTASE (GLUCOSE PHOSPHOMUTASE) (PGM)	od55d04.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1371847.3*	601110258F1 NIH_MGC_16 Hamo sapiens cDNA clone IMAGE:3350750 5'	601110258F1 NIH_MGC_16 Hamo sapiens cDNA clane IMAGE:3350750 5'	0x42c10.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE:2272242 3'	Homo sapiens X28 region near ALD locus containing dual specificity phosphatase 9 (DUSP9), ribosomal	protein L18a (RPL18a), Ca2+/Calmodulin-dependent protein kinase I (CAMKI), creatine transporter (CRTR),	CDM protein (CDM), adrendeukodystrophy protein >	Drosophila melanogaster sodium channel protein (para) gene, exons 9,10,11,12 and optional segments b, c, d	and e, partial cds	Triticum aestivum stripe rust resistance protein Yr10 (Yr10) gene, complete cds	Salmonella typhimurium adenine-methytransferase (mod) and restriction endonuclease (res)
Exon Probes E	Top Hit Database Source	EST_HUMAN E	O LN		H IN	NT TN	SWISSPROT	ISSPROT	NT IN	SWISSPROT B	o LN	SWISSPROT A	O IN	X X	SWISSPROT	Г	П	3		W ≅ E		EST_HUMAN 6	SWISSPROT P				EST_HUMAN D	Ξ.		NT				NT S
Single	Top Hit Acession No.	AW976184.1	AF245455.1	AF245455.1	AL163302.2	AF174585.1	P49657	Q09632	U65667.1	Q28642	AJ005029.1	P22567	AJ003108.1	AF174844.1	067551	BE957439.2	BE957439.2	A 12024EB 4	73302130.1	AJ302158.1	BF034016.1	BF034016.1	P38652	AA825565.1	BE258705.1	BE258705.1	A1680876.1			U52111.2		U26716.1	AF149112.1	9.7E-01 M90544.1
	Most Similar (Top) Hit BLAST E Value	1.0E+00	9.9E-01	9.9E-01	9.9E-01		9.9E-01			9.9E-01			9.8E-01				9.8E-01	20 00	9.00	9.8E-01	9.8E-01		9.8E-01		9.8E-01	9.8E-01				9.8E-01		9.7E-01	9.7E-01	9.7E-01
	Expression Signal	3.08	16.0	0.97	1.17	96.0	14.59	0.83	1.39	2.61	1.08	1.77	0.89	2.05	0.95	0.61	0.61	30 7	8	4.86	1.13	1.13	7.0	0.56	4.86	4.86	1.78			1.39		2.51	1.7	1.28
	ORF SEQ ID NO:	-	28742	26743	27794		31162	31389			36142	25658			28930	28933	28934		*2075	32635	82028	62028	34110				37109							33901
Ī	Exon SEQ ID NO:	24410	14209	14209	15222	16267	18440		21702	21873	23128	13180	14907	15379	16467	16470	18470	02207	1	19779					23395	23395	24040			24341		- 1		20986
	Probe SEQ ID NO:	12171	1616	1616	2664	3665	.5816	6029	9185	9474	10593	549	2336	2827	3869	3872	3872	7750	37,	7250	7641	7641	8653	10334	10874	10874	11597			12058		7212	8440	8448

Page 29 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

_		т.	_	_	_	_	_	_		_			_	_	_	_				-	_	_	_	_	_	_	T .	_		.		_
	Top Hit Descriptor	UI-H-BI4-aci-e-07-0-UI.s1 NCI_CGAP_Sub8 Homo sapiens cDNA clone IMACE:3085140 3'	Botrytis cinerea strain T4 cDNA library under conditions of nitrogen deprivation	Bromus inermis putative cytosolic phosphoglucomutase (pgm1) mRNA, complete cds	Bromus inermis putative cytosolic phosphoglucomutase (pgm1) mRNA, complete cds	PM2-UM0053-240300-005-f12 UM0053 Homo sepiens cDNA	Panovirus B19 DNA, patient C, genome position 2448-2994	Parvovirus B19 DNA, patient C, genome position 2448-2994	P. falciparum complete gene map of plastid-like DNA (IR-A)	Rattus norvegicus (strain R21) Rps2r gene, complete cds	Mus musculus WNT-2 gene, partial cds; putative ankyrin-related protein and cystic fibrosis transmembrane	Conductation regulated (LPTIX) genes, section 1 of 2 of the complete cost, and unknown gene Homo segions abosomal protein s.4 Visitions name complete cite	AV752805 NPD Home senions of the Alma NPDRACIAS 5	AV752605 NPD Homo sapiens cDNA clone NPDBAG06 5'	Homo saplens centrosomal protein 2 (CEP2), mRNA	Sphyrna tiburo NADH dehydrogenase subunit 2 (NADH2) gene, mitochondrial gene encoding mitochondrial	protein, partial cds	Homo sapiens CGI-125 protein (LOC51003), mRNA	ENDOGLUCANASE I PRECURSOR (EGI) (ENDO-1,4-BETA-GLUCANASE) (CELLULASE I)	601675639F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3958473 5	601675639F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3958473 5'	qd57d07.x1 Soares_tests_NHT Homo sepiens cDNA clone IMAGE:1733581 3'	RC1-C10295-241199-011-b02 C10295 Homo sapiens cDNA	801885163F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4103630 5'	UI-H-BI2-ahp-f-03-0-UI.s1 NCI_CGAP_Sub4 Homo sapiens cDNA clone IMAGE:2727677 3'	Bartonella clarridgetae RNA polymerase beta subunit (rpoB) gene, partial cds	Pimpinella brachycarpa zinc finger protein (ZFP1) mRNA, complete cds	Human Fc-gamma-receptorIIA (FCGR2A) gene, exon 4	Нотто sapiens phytanoyl-CoA hydroxylase (PHYH) gene, exon 5	RC5-BT0503-271199-011-B01 BT0503 Homo sapiens cDNA	Bovine papillomavirus type 2, complete genome	Bovine papillomavirus type 2, complete genome
	Top Hit Database Source	EST_HUMAN	LN	N	N	EST_HUMAN	۲	LN L	LN L	μ	1	Z	FOT HIMAN	EST HUMAN	NT.		N⊤	LN	SWISSPROT	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	NT	LN	IN	LN	EST_HUMAN	NT	NT
	Top Hit Acession No.	BF511209.1	AL114281.1	AF197925.1	AF197925.1	AW 799674.1	Z70556.1	Z70556.1	X85275.1	L81138.1	* 6*00000	AF041427 1	AV752805 1	AV752805.1	11421722 NT		U91423.1	7705591 NT	002834	BE902340.1	BE902340.1	Al190162.1	AW861102.1	BF218771.1	AW 293799.1	AF165990.1	AF080595.1	M90724.1	AF242382.1	BE071172.1	M20219.1	1 M20219.1
	Most Similar (Top) Hit BLAST E Value	9.7E-01	9.7E-01	9.6E-01	9.6E-01		_	9.6E-01	9.6E-01	9.6E-01		9.0E-01		9.6E-01	9.6E-01		=	9.5E-01	9.5E-01	9.5E-01		9.5E-01 /	9.5E-01	9.5E-01	9.5E-01	9.4E-01	9.4E-01	9.4E-01	9.3E-01	9.3E-01	9.3E-01	9.3E-01
	Expression Signal	5.23	2.92	0.58	0.58	1.71	3.9	3.9	1.23	0.47		181	5 18	5.18	2.38		2.8	1.02	1.2	1.89	1.89	0.63	1.07	1.71	1.59	1.8	2.47	0.88	0.85	1.09	0.92	0.92
	ORF SEQ ID NO:			59559	29560			31277		34248	30000	38507	36912	36913					27817	28909	28910	34387	34500	38674	38033			34265		27792	29154	29155
	Exan SEQ ID NO:	23562		17115	17115		18550		20872			1	23847	1.	1		24983			16448	16448	21469	21571	23831	23024	15847		21338	14358		16701	16701
	Probe SEQ ID NO:	11049	12658	4531	4531	4557	5928	5928	8331	8785	000	10966	11395	11395	11733		12388	2515	2691	3850	3850	8931	9034	11123	11326	3235	3254	8799	1768	2662	4107	4107

Page 30 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Probe	Exon	ORF SEQ	Expression	Most Similar (Top) Hit	Top Hit Acession	Top Hit	Too Hit Descriptor
S S	S S	Ö NÖ:	Signal	BLAST E Value	ó Z	Source	. Tradisco de la constante de
5778	18403	31119	1.41	9.3E-01	AF213884.1	Ņ	Homo saplens nuclear factor of kappa light polypeptide gene enhancer in B-cells 1 (NFKB1) gene, complete ods
5858	L	31204		9.3E-01	L36189.1	F	Spodoptera frugiperda methylenetatrahydrofolata dehydrogenase mRNA, complete cds
8011	20553		1.62	9.3E-01	AA847040.1	EST_HUMAN	oe09b03.s1 NCI_CGAP_Ov2 Hamo sapiens dDNA clone IMAGE:1385357
8748			1.13		AF061981.1	NT	Xenopus laevis CCCH zinc finger protein C3H-2 (C3H-2) mRNA, complete cds
8867		34330	1.01	9.3E-01	AL161534.2	NT	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 34
12508	24629		1.87	9.3E-01	11440298 NT	TN	Homo sapiens inositol 1,4,5-triphosphate receptor, type 2 (ITPR2), mRNA
12515	24634		2	9.3E-01	AF271207.1	IN	Aedes triseriatus putative large subunit ribosomal protein rpL34 mRNA, complete cds
3276	15887	28369	3.99	9.2E-01	BE622702.1	EST_HUMAN	601441338T1 NIH_MGC_72 Hamo sapiens cDNA clone IMAGE:3916184 3'
5004	17577		0.62		BF129973.1	EST_HUMAN	601817814F1 NIH_MGC_58 Homo sapiens cDNA clone IMAGE:4041363 5'
5894	18516		1,41	9.2E-01	7106410 NT	NT	Mus musculus solute carrier family 30 (zinc transporter), member 4 (Slc30a4), mRNA
6140	18754		4.4		BF037586.1	EST_HUMAN	801461153F1 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:3864661 5'
9278	22078	35042	1.31	9.2E-01	AL161565.2	TN.	Arabidopsis thallana DNA chromosome 4, contig fragment No. 65
9663	22182	35135	1.15	9.2E-01	6671677 NT	NT	Mus musculus carbonic anhydrase 4 (Car4), mRNA
10166	22661	35656	3.47	9.2E-01	11430963 NT	NT	Homo sapiens lysosomal apyrase-like protein 1 (LALP1), mRNA
10314	22808	35800	1.58	9.2E-01	BF593251.1	EST HUMAN	7o58e08.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:3578219 3' similar to SW:NU5M_TRYBB P04540 NADH-UBIQUINONE OXIDOREDUCTASE CHAIN 5 :
10526			:			EST_HUMAN	601334943F1 NIH_MGC_39 Homo sapiens cDNA clone IMAGE:3688714 5'
11569	24016				BF132402.1	EST_HUMAN	601820312F1 NIH_MGC_58 Hamo sapiens cDNA clone IMAGE:4052018 5'
1666	14259	28793	4 89		196675 1	EST HUMAN	ye52f01.s1 Soares fetal liver spleen 1NFLS Homo sepiens cDNA clone IMAGE:121369 3' similar to contains. Alu recettiive element:
2169	1					LN	Homo sapiens hypothetical protein FLJ20048 (FLJ20048), mRNA
3239	15851	28331	0.83	9.1E-01	T26418.1	EST_HUMAN	AB200GBR Infant brain, LLNL array of Dr. M. Soares 1NIB Homo sapiens cDNA done LLAB200G8 5
3239	15851	28332	0.83	9.1E-01	T26418.1	EST HUMAN	AB200G8R Infant brain, LLNL array of Dr. M. Soares 1NIB Homo sapiens cDNA clone LLAB200G8 5'
6315					L36033.1	N	Human pre-B cell stimulating factor homologue (SDF1b) mRNA, complete cds
9630	19226		2.82	9.1E-01	Q61704	SWISSPROT	INTER-ALPHA-TRYPSIN INHIBITOR HEAVY CHAIN H3 PRECURSOR (ITI HEAVY CHAIN H3)
7577	20093	32970	15.95	9.1E-01	AA806623.1	EST_HUMAN	ob71g08.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1338862 3'
7719					U72995.1	LN	Rattus norvegicus Rab3 GDP/GTP exchange protain mRNA, complete cds
12093			33.14		AF05011	N	Homo sapiens uncoupling protein-3 (UCP3) gene, complete cds
3241		28335		·		LΝ	Homo saplens DKFZP564M2423 protein (DKFZP584M2423), mRNA
3401		`.				IN	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 27
4468	17054	29498	1.44	9.0E-01	AF099810.1	NT	Homo sapiens neuredn III-alpha gene, partial ods

Page 31 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Danio rerio LIM class homeodomain protein (lim5) mRNA, complete cds	Xenopus laevis gene for aldolase, complete cds	Danio rerio semaphorin Z1a mRNA, complete cds	Mycoplasma genitalium section 24 of 51 of the complete genome	Fugu rubripes neural cell adhesion molecule L1 homolog (L1-CAM) gene, complete (xis; putative protein 1 (P) IT1) neura narial cris: mitreles perfilicity chromosome sententin protein SMC1 by noton (SMC1) neura	complete cds; and calcium channel apha-1 subunit>	Rabbit MHC fragment RLA-DF DNA	Homo sapiens TESTIN 2 and TESTIN 3 genes, complete cds, alternatively spliced	Oithona nana cytochrome-c oxidase subunit I (coxl) gene, partial cds; mitochondrial gene for mitochondrial	Xylella fastidiosa, section 90 of 229 of the complete genome	Chlamydophlia pneumontae AR39, section 21 of 94 of the complete genome	Т	Pseudorabies virus Ea glycoprotein M gene, complete cds	M.eeruginosa (HUB 5-2-4) DNA from plasmid PMA1	Synechocystis sp. PCC8803 complete genome, 13/27, 1578593-1719643	Homo saplens SOS1 (SOS1) gene, partial cds	Homo sapiens AT-binding transcription factor 1 (ATBF1), mRNA	nn05/11.s1 NCI_CGAP_P14.1 Homo sapiens cDNA clone IMAGE:1076877	Homo sapiens xeroderma pigmentosum complementation group C (XPC) gene, intron 9	Homo sapiens xeroderma pigmentosum complementation group C (XPC) gene, intron 9	Pseudomonas aeruginosa topoisomerase (top), putative transcriptional regulatory protein OhbR (chbR), ortho-	Havodatozate 1,z-ciroxygenese beta-157 protein Orba (orba), Orbo (orbo), d'indriandenzate 1,z- Idioxogenase alphe-18P protein Obb8 (obb8), and out?		П	Г	Pseudomonas aeruginosa PA01, section 524 of 529 of the complete genome			П		601823684R1 NIH_MGC_79 Homo sepiens cDNA clone IMAGE:4043564 3'
Top Hit Database Source	TN	NT	NT	TN		F	TN	N	Į.	Į	Z	SWISSPROT	Z	TN	NT	NT	TN	EST_HUMAN	IN	NT		_ <u>_</u>	EST HUMAN	EST_HUMAN	EST_HUMAN	LΝ	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EŞT_HUMAN
Top Hit Acession No.	9.0E-01 L42547.1	11 D38621.1	9.0E-01 AF086761.1	9.0E-01 U39702.1		AF026198.1	8.9E-01 X60986.1	8.9E-01 AF260225.1	8 OE -01 A F2550687 1	8 9F-01 AF003944 1	8.9E-01 AE002186.2	026350	AF310617.1	8.8E-01 Z28337.1	8.8E-01 D90911.1	8.7E-01 AF106953.2	5901893 NT	8.7E-01 AA595863.1	1 AF156539.1	8.7E-01 AF156539.1		8 7E-01 AF121970.1	1 AW897335.1	1 AI239456.1	11 AI239456.1	1 AE004963.1	1 BF570169.1	1 BF570169.1	8.7E-01 BF363970.1	1 BF107694.1	01 BF107694.1
Most Similar (Top) Hit BLAST E Value	9.0E-01	9.0E-01	9.0E-01	9.0E-01		8.9E-01	8.9E-01	8.9E-01	9 OF 01	8 9F-01	8.9E-01	8.8E-01 026350	8.8E-01	8.8E-01	8.8E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01		8 7F-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01	8.7E-01
Expression Signal	0.78	1.64	0.54	0.47		2.49	1.27	0.47		2 59	5.33	2.1	0.7	3.82	2.27	1.48	1,13	5.67	0.61	0.61		308	0.66	0.75	0.75	1.7	95.0	0.56	5.79	4.31	4.31
ORF SEQ ID NO:	32814		34746			31222			70866		L	L	30658			25609	27585			29876			33431		34323	35122	35693				37098
Exan SEQ ID NO:			21797	22242		18497	L	20693	SUBUC				L		25067	13123	15013		17423	17423		17721	1_	21399	L		<u> </u>			L	24028
Probe SEQ ID NO:	7424	7450	9271	9744		5875	6396	8152	9385	11818	11927	4840	5576	10960	11749	490	2446	2898	4845	4845		5151	7983	988	9860	9653	10205	10205	10711	11582	11582

Page 32 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	AV661898 GLC Homo sapiens cDNA clone GLCGYG073'	Rat iGFII gene for insulin-like growth factor II	zd44e03.r1 Soares_fetal_heart_NbHH19W Homo sapiens cDNA clone IMAGE:343516 5	Homo sapiens cytochrome P450, subfamily XXVIIA (steroid 27-hydroxylase, cerebrotendinous xambomatosis), polyoeptide 1 (CYP27A1b) mRNA	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 65	Drosophila melanogaster merlin (Dmerlin) mRNA, complete cds	Chicken lipoprotein lipase gene	Chicken lipoprotein lipase gene	Grus canadensis recombination activating protein 1 (RAG-1) gene, partial cds	Grus canadensis recombination activating protein 1 (RAG-1) gene, partial cds	Bacilius halodurans genomic DNA, section 12/14	Drosophila melanogaster collapsin response mediator protein (CRMP) mRNA, complete cds	Archaeoglobus fulgidus section 128 of 172 of the complete genome	Botrytis cinerea strain T4 cDNA library under conditions of nitrogen deprivation	Bacteriophage D3, complete genome	801087107F1 NIH_MGC_10 Hamo sapiens cDNA clane IMAGE:3453505 5'	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 68	SEGMENTATION PROTEIN PAIRED	SEGMENTATION PROTEIN PAIRED	Homo sapiens partial 54T4 receptor gene, exons 2 to 5	Cyanidium caldarium gene for SigC, complete cds	Cyanidium caldarium gene for SigC, complete cds	Homo sapiens human immunodeficiency virus type I enhancer-binding protein 1 (HIVEP1), mRNA	Rattus norvegicus protein tyrosine phosphatase, non-receptor type 5 (Ptpn5), mRNA	Fowl adenovirus 8, complete genome	Human fibroblast growth factor receptor 3 (FGFR3) gene, intron 7	Human fibroblast growth factor receptor 3 (FGFR3) gene, intron 7	Pyrococcus abyssi complete genome; segment 5/6	Thermus thermophilus cytochrome c-552 (cycA) and CycB (cycB) genes, complete cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 18	Nicotiana tabacum mRNA for chloroplast ribosomal protein L10, complete cds	Streptomyces antibioticus polyketide biosynthetic gene cluster	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 40
Top Hit Database Source	EST_HUMAN A	TN PA	EST_HUMAN 2					TN TN	Ę	IN		IN	/ IN		NT TN	T_HUMAN		SWISSPROT	SWISSPROT	LN ⊢N	LN	LN			I LN	LN	N N	LN LN		TN.		IN	
Top Hit Acession No.	AV661898.1	X17012.1	_	TN 01-0503-0	AL161565.2			X60547.1	AF143732.1	AF143732.1	AP001518.1	AF077837.1	AE000979.1	AL112162.1	AF165214.1	BE542612.1	AL161572.2	P06601	P06601	AJ243213.1	AB006799.1	AB006799.1	11418543 NT	9507008 NT	AF083975.2	L78726.1	L78726.1	AJ248287.1	M93437.1	AL161506.2	AB010879.1	Y19177.1	AL161540.2
Most Similar (Top) Hit BLAST E Vatue	8.7E-01	8.6E-01	8.6E-01	8 6F-01	-		+	8.6E-01	8.6E-01	8.6E-01	8.6E-01	8.6E-01	8.6E-01	8.6E-01	8.5E-01			8.5E-01	8.5E-01	8.5E-01	8.5E-01	8.5E-01	8.5E-01	8.5E-01	8.4E-01	8.4E-01	8.4E-01	8.4E-01	8.3E-01	8.3E-01	8.3E-01		8.3E-01
Expression Signal	4.44	1.55	8.72	108	0.78	1.38	90.6	90'6	1.88	1.88	1.33	23.0	0.48	1.73	1.32	2.38	15.0	0.84	0.84	0.51	1.38	1.38	3.12	7.92	0.62	3.15	3.15	2.68	2.48	3.26	99.0	3.24	2.15
ORF SEQ ID NO:			26024	27457		28931		31416	32216	32217		33434			32232	32928	33383	33817	33818	33896	35734	35735			29890	30807	30808	l 	25889	28212			Ш
Exen SEQ ID NO:	24861	13132	.13505			16468	18674	18674	19401	19401	20410	20528	22103	24812	19416	20053	20474	20897		20981	22747	22747	24978	24355	17440	24747	24747	22365	13390	L	16481	16880	18107
Probe SEQ ID NO:	12146	တ္တ	894	23.10	3681	3870	6057	6057	6810	6810	7868	7986	9803	12338	6826	7533	7932	8357	8357	8441	10252	10252	12077	12084	4862	2895	5685	9986	771	3129	3883	4084	5473

Page 33 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

ſ		£	Γ	Γ		Ť		Γ				Γ		Γ			_	Γ									T	T		7	
	Top Hit Descriptor	nn01112.y5 NCI_CGAP_Co9 Homo sapiens cDNA clone IMAGE:1076495 5' simitar to contains THR.t1 THR repetitive element ;	Drosophila melanogaster Lis 1 homolog mRNA, complete cds	Mus musculus neuro-d4 gene, exons 3 through 12 and partial cds	Methanobacterium thermoautotrophicum from bases 1270510 to 1283409 (section 109 of 148) of the complete genome	Phytophthora infestans mitochondrion, complete genome	Homo sapiens FRA3B common fragile region, diadenosine triphosphate hydrolase (FHIT) gene, exon 5	Rattus norvegicus mRNA for RPHO-1, complete cds	Mus musculus trophinin (Tnn) gene, complete cds	Homo sapiens mRNA for KIAA0674 protein, partial cds	S. cerevisiae chromosome VII reading frame ORF YGL062w	S. cerevisiae chromosome VII reading frame ORF YGL062w	Rattus norvegicus mRNA for RPHO-1, complete cds	Homo sapiens mRNA for KIAA 1034 protein, partial cds	Amanita muscaria mRNA for SCIII25 protein	CM4-HT0243-081189-037-e01 HT0243 Homo sapiens cDNA	S.cerevisiae MET, LEU4, and POL1 genes encoding MET4 protein, alpha-isoproplymalate (alpha-IPM) synthetase (partial), and DNA polymerase alpha (partial)	Homo sapiens mRNA for KIAA0630 protein, partial cds	Homo sapiens thioredoxin-related protein mRNA, complete cds	Oncorhynchus Ishawytscha isolate T-20 somatolactin precursor gene, exon 1	Oncorhynchus tshawytscha isolate T-20 somatolactin precursor gene, exon 1	MCKUSICK-KAUFMAN/BARDET-BIEDL SYNDROMES PUTATIVE CHAPERONIN	MCKUSICK-KAUFMAN/BARDET-BIEDL SYNDROMES PUTATIVE CHAPERONIN	Moltuscum contegiosum virus type 1 ORF1 and ORF2 DNA	OVARIAN TUMOR LOCUS PROTEIN	yw14d02.r1 Soares_placenta_8to9weeks_2NbHP8to9W Home sapiens cDNA clone IMAGE:252195 5'	SILILIAN TO BUINDON'S GOOD MIDOURADE THO (TOWNIN).	Mus musculus mKNA tor NIPSNAPZ protein	Mus musculus TANK binding kinase TBK1 (Tbk1) mRNA, complete cds	Homo sapiens MHC class 1 region	Homo sapiens MHC class 1 region
	Top Hit Database Source	EST_HUMAN	Ę	ΙN	ĻΝ	Z	Z	TN	NT	NT	LN	FZ	LN	LN LN	LZ	EST_HUMAN	LN	LΝ	LX	L	IN	SWISSPROT	SWISSPROT	LN	SWISSPROT	144941 111 200	NAMOR I SE	LN	Ā	Ŋ	뉟
	Top Hit Acession No.	AI791952.1	AF098070.1	AF108133.1	AE000903.1	2472	AF020503.1	AB000489.1	AF145589.1	AB014574.1	272584.1	Z72584.1	AB000489.1	AB028957.1	AJ010142.1	AW379433.1	212126.1	AB014530.1	AF052859.1	AF223888.1	AF223888.1	021170	1 09J170	L10127.1	P10383	, 000201	8.ZE-U1 F187396.1	1 AJ001261.1	8.1E-01 AF191839.1	8.1E-01 AF055068.1	8.1E-01 AF055066.1
	Most Similar (Top) Hit BLAST E Value		_	8.3E-01	8.3E-01	_	8.3E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01		8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	8.2E-01	_	8.2E-01	100	0.42-01	8.2E-01	8.1E-01	8.1E-01	8.1E-01
	Expression Signal	3.14	1,11	3.5	2.92	2.52	2.45	3.23	1.45	1.12	0.61	0.61	1.08	2.11	8.0	3.18	4.21	0.63	1.67	0.59	0.59	3.52	3.52	3.33	8.05		8 9	1.86	1.79	2.99	2.89
	ORF SEQ ID NO:		35507	35604	36103		36735	27244		28038				30385	32439	32379	32700	L	35450		35610	35772	35773	37017	37091			30970			28586
	Exon SEQ ID NO:	22087	l	22614	l		23688	14675	14715	16567	ı	16798		17971	19605	19554	24779			22618			Ĺ	L	L		1		15340		16109
	Probe SEQ ID NO:	9587	10019	10119	10553	10571	11183	2096	2137	3969	4209	4209	5270	5420	6871	8269	7313	9638	9971	10123	10123	10286	10286	11498	11576	1	3	12102	2787	3504	3504

Page 34 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

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Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Simitar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
5046	17619		99.0	8.1E-01	4.1	LN	Drosophila melanogaster Na/K-ATPase beta subunit isoform 4 (JYbeta2) mRNA, complete cds
6457	19058		88.0	8.1E-01	U16790.1	. LN	Mus musculus putative collagen alpha-2 (XI) chain (COL11A2) gene, partial cds
6735	19329			8.1E-01	Q13491		NEURONAL MEMBRANE GLYCOPROTEIN M6-B
6735	19329		2.54	8.1E-01	Q13491		NEURONAL MEMBRANE GLYCOPROTEIN M6-B
7853	70304	90000	70 0	n v	A E000749 0	£1	Drosophila melanogaster putative inorganic phosphate cotransporter (Picot) gene, partial cds; putative sodium channel (Nach) and putative amylase-related protein (Amyrel) genes, complete cds; and putative serine-
扌	2000						and form of the second control of the second
							Drosophila melanogaster putative inorganic phosphate cotransporter (Picot) gene, partial cds, putative sodium channel (Nach) and putative amylase-related protein (Amyrel) genes, complete cds; and putative serine-
7852	20394				AF022713.2	۲	enriched protein (gprs) gene, partial cd>
8545	21084		0.92	8.1E-01	AP001517.1		Bacillus halodurans genomic DNA, section 11/14
8545	21084	34007		8.1E-01	AP001517.1	INI	Bacillus halodurans genomic DNA, section 11/14
							xn01h03.x1 NCI_CGAP_Kid11 Hamo septens cDNA clone IMAGE:2892469 3' similar to SW:LYAR_MOUSE Q08288 CELL GROWTH REGULATING NUCLEOLAR PROTEIN; contains MER22.b1 PTR5 repetitive
8705	21244	34167	1.08	8.1E-01	AW242647.1	EST_HUMAN	element;
10032	22527			8.1E-01	P06425	SWISSPROT	PROBABLE E4 PROTEIN
11356	23810		2.97	8.1E-01	BE838558.1	EST_HUMAN	RC0-TN0080-220800-025-410 TN0080 Homo sapiens cDNA
11356	23810					EST_HUMAN	RC0-TN0080-220800-025-d10 TN0080 Homo sapiens cDNA
11811	24183	31031			AE001711.1		Thermotoga maritima section 23 of 136 of the complete genome
188	12849		4.99	8.0E-01	AJ271510.1		Staphylococcus aureus partial pta gene for phosphate actyltransferase allele 15
310	12985	25453	7.95	8.0E-01	AJ132772.1	TN	Bos taurus futb and rtif genes
2080	14661				BF530962.1	THUMAN	602072473F1 NCI_CGAP_Brn67 Homo sapiens cDNA clone IMAGE:4215091 5'
3113	15728		1.24	8.0E-01	AF127897.1		Saimin boliviensis offactory receptor (SBO27) gene, partial cds
3354	15962		1.13	8.0E-01	AB006193.1	LΝ	Mus musculus gene for oxiductal glycoprotein, complete cds
3765	16366		1.05	8.0E-01	AL162758.2	IN	Neisseria meningitidis serogroup A strain Z2491 complete genome; segment 7/7
4	17213	29664			X83739.2	N T	G.gallus mRNA for nicotinic acetylcholine receptor (nAChR) beta 3 subunit
5117	17689		1.09	8.0E-01	7657352 NT		Mus musculus myosin IXb (Myo9b), mRNA
188	20473		2:32	8.0E-01		EST_HUMAN	RC0-NN1012-270300-021-h06 NN1012 Homo sapiens cDNA
8462	21002	33919	1.17	8.0E-01	Y11095.1	NT	Rice stripe virus RNA 3
479	13112	25602	1.37	7.9E-01	D11476.1	μ	Lymantria dispar nuclear polyhedrosis virus gene for DNA polymerase, complete cds
744	13364		1.05	7.9E-01	AE002130.1	NT.	Ureaplasma urealyticum section 31 of 59 of the complete genome
1648	14240		28.9		AB040885.1	NT	Homo sapiens mRNA for KIAA1452 protein, partial cds
1695	14288		1.11		U32739.1	NT	Haemophilus influenzae Rd section 54 of 163 of the complete genome
2303	14878	27452	8.76	7.9E-01	AB004816.1	LN	Oryctolagus cuniculus mRNA for mitsugumin 29, complete cds
		İ					

Page 35 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 36 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Detabase Source	_			
	ġ			(Top) Hit BLAST E Value
38/NT (GALNAC-17) (GALNAC-17), MKNA	8393408 NT	7.7E-01 839340		7.7E-01
NT Homo sapiens PRO1975 mRNA, complete cds	F118085.1	7.7E-01 AF118085.1		7.7E-01
	F199488.1	7.7E-01 AF199488.1		3.17 7.7E-01
NT Cotumix cotumix japonica sub-species japonica beta-actin mRNA, partial cds	F199488.1	7.7E-01 AF199488.1		7.7E-01
SWISSPROT RAFFINOSE INVERTASE (INVERTASE)	16553	7.7E-01 P16553		7.7E-01
SWISSPROT RAFFINOSE INVERTASE (INVERTASE)		P16553	P16553	7.7E-01 P16553
EST_HUMAN yf24b02.s1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:127755 3'		R08600.1	R08600.1	0.8] 7.7E-01]R08600.1
NT Daphnia magna hemoglobin gene cluster (dhb3, dhb1 and dhb2 genes), complete cds		AB021134.1	AB021134.1	7.7E-01 AB021134.1
T Archaeoglobus fulgidus, complete genome	11497621 NT	7.7E-01 11497621		7.7E-01
NT Overlogus curiculus immaglobulin VDV region gene		127316.1	127316.1	7.6E-01 L27316.1
NT Oryctolagus cuniculus immunoglobulin VDJ region gene		L27316.1	L27316.1	19.73 7.6E-01 [L27316.1
Arabidopsis thaliana 3-methylcrotonyl-CoA carboxylase non-biotinylated subunit (MCCB) mRNA, complete cds		AF059510.1	AF059510.1	7.6E-01 AF059510.1
NT cds Cds Cdirector MATING TYPE DBOTEIN & AI DH& 74	10.1	7.6E-01 AF059510.1	AF059510.1	7.6E-01 AF059610.1
T	T	A 1252200 4	7.05 4.05 4.05	0.00 1.00-0.1
EST HUMAN (add 140 Z.X.) Stanlay Frontal NS pool 2 Home sapiens convectione link GF 2030879		7 AE 01 A1253388	7 85-01 412533393	0.850 7.0E-10.1A12533383.1
		7.8E-01 U72487.1	7.8E-01 U72487.1	0.98 7.8E-01 U72487.1
Mus musculus neuromedin U precursor (Nmu) gene, partial cds; tPhLP (Tphlp) gene, partial cds; CLOCK (Clock) gene, complete cds; PFT27 (Pft27) gene, complete cds; and H5AR (H5ar) gene, complete cds		7.6E-01 AF146793.2 N		7.6E-01 AF146793.2
	6857752 NT			1.76 7.6E-01
	. 6857752 NT	7.6E-01 . 6857752 N		7.6E-01
GLUTAMATE INMDA RECEPTOR SUBUNIT EPSILON 3 PRECURSOR (N-METHYL D-ASPARTATE				
SWISSPROT RECEPTOR		7.6E-01 C01098		7.6E-01 C01098
GLUTAMATE [NMDA] RECEPTOR SUBUNIT EPSILON 3 PRECURSOR (N-METHYL D-ASPARTATE SWISSPROT RECEPTOR SUBTYPE 2C) (NR2C) (NMDAR2C)		7.6E-01 Q01098		7.6E-01 Q01098
T Mus musculus cytochrome P450, 2b9, phenobarbitol inducible, type a (Cyp2b9), mRNA	6753577 NT	7.6E-01 6753577 N		7.6E-01
SWISSPROT MUSCARINIC ACETYLCHOLINE RECEPTOR M2	30372			3.33
SWISSPROT MUSCARINIC ACETYLCHOLINE RECEPTOR M2	30372	7.6E-01 P30372	3.33 7.6E-01 P30372	
	86347.1			2.74
NT H. aspersa mRNA for neurofilament NF70	86347.1		2.74 7.6E-01 X86347.1	

Page 37 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Porber SEO Exm Port SEO Port SEO Port SEO Top HILD Descriptor Top HILD	+							
24004 5.74 7.6E-01 AL161592.2 NT 24121 6.31 7.6E-01 AL161592.2 NT 13170 1.32 7.5E-01 AL163301.2 NT 24318 25712 1.13 7.5E-01 AL163301.2 NT 24318 25723 0.74 7.5E-01 AF16331.2 NT 24318 30897 1.31 7.5E-01 AF16331.2 NT 24318 30897 1.31 7.5E-01 AF162381.1 NT 24318 30897 1.31 7.4E-01 AF132381.1 NT 24328 30897 1.31 7.4E-01 AF132381.1 NT 24328 30897 1.31 7.4E-01 AF132381.1 NT 20328 33236 1.03 7.4E-01 AF185346.1 EST_HUMAN 21107 34028 0.69 7.4E-01 AF185346.1 NT 21107 34028 0.69 7.4E-01 BE745626.1 NT 21107 </td <td></td> <td>Exan SEQ ID NO:</td> <td>ORF SEQ ID NO:</td> <td>Expression Signal</td> <td>Most Similar (Top) Hit BLAST E Value</td> <td>Top Hit Acession No.</td> <td>Top Hit Database Source</td> <td>Top Hit Descriptor</td>		Exan SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
24121 6.31 7.6E-01 AB020702.1 NT 13128 25712 1.32 7.5E-01 AL163301.2 NT 20050 32923 0.74 7.5E-01 AF020503.1 NT 20050 32923 0.74 7.5E-01 AF052730.1 NT 24318 30897 1.91 7.5E-01 AF052730.1 NT 24638 30897 1.91 7.5E-01 AF052730.1 NT 13771 26279 1.36 7.4E-01 AF16334.2 NT 13771 26279 1.36 7.4E-01 AF16324.2 NT 13771 26279 1.37 7.4E-01 AF16324.2 NT 20328 33234 1.03 7.4E-01 AF16532.2 NT 21107 34028 0.63 7.4E-01 BF74526.1 RST HUMAN 21107 34028 0.64 7.4E-01 BF74503.1 EST HUMAN 21107 3458 1.19 7.4E-01 AA187286.1 <t< td=""><td></td><td>24004</td><td></td><td>5.74</td><td>7.6E-01</td><td>П</td><td>NT</td><td>Arabidopsis thaliana DNA chromosome 4, contig fregment No. 88</td></t<>		24004		5.74	7.6E-01	П	NT	Arabidopsis thaliana DNA chromosome 4, contig fregment No. 88
132 7.5E-01 AL163301.2 NT 20056 32923 0.74 7.5E-01 AF020303.1 NT 24318 25712 1.13 7.5E-01 AF052730.1 NT 24318 32923 0.74 7.5E-01 AF052730.1 NT 24638 30697 1.31 7.5E-01 AF052730.1 NT 13771 28278 1.36 7.4E-01 AF112538.1 NT 16386 28430 7.7 7.4E-01 AF112538.1 NT 20328 33224 1.03 7.4E-01 AL16351.2 NT 21107 34026 0.64 7.4E-01 AL16351.2 NT 21107 34026 0.69 7.4E-01 AL161551.2 NT 21107 34026 0.69 7.4E-01 AL161561.2 NT 21107 34026 0.69 7.4E-01 AR161661.2 NT 21107 34026 0.69 7.4E-01 AR161660.1 NT	Щ	24121		6.31			NT	Homo sapiens mRNA for KIAA0895 protein, partial cds
13238 25712 1.13 7.5E-01 AF052730.1 NT 20050 32823 0.74 7.5E-01 AF052730.1 NT 24318 30897 1.91 7.5E-01 AF052730.1 NT 24318 30897 1.91 7.5E-01 AF052730.1 NT 13771 28279 1.36 7.4E-01 AF12538.1 NT 16385 28430 7.7 7.4E-01 AF12538.1 NT 20328 33234 1.03 7.4E-01 AL163246.2 NT 20328 33235 1.03 7.4E-01 AL163246.2 NT 21107 34026 0.69 7.4E-01 BE747503.1 EST_HUMAN 21563 34492 7.17 7.4E-01 BE747503.1 EST_HUMAN 21649 37034 1.68 7.4E-01 BE747503.1 EST_HUMAN 22705 35786 0.59 7.4E-01 AR17869.2 NT 23984 37034 1.68 7.4E-01 AR17869.2 NT 24076 35786 0.59 7.4E-01 AR17869.1 NT 24078<	Щ	13170		1.32	_	П	NT	Homo sapiens chronosome 21 segment HS21C101
20050 32923 0.74 7.5E-01 AF163151.2 NT 24318 5.28 7.5E-01 AF163151.2 NT 24318 5.29 7.5E-01 D90907.1 NT 13771 24318 7.4E-01 AF16338.1 NT 16389 28854 0.63 7.4E-01 AF16338.1 NT 20328 33234 1.03 7.4E-01 AL163248.2 NT 20328 33234 1.03 7.4E-01 AL161551.2 NT 20328 33234 1.03 7.4E-01 AL161551.2 NT 21107 34026 0.64 7.4E-01 BF34626.1 EST_HUMAN 21186 34492 7.17 7.4E-01 BF34626.1 BST_HUMAN 21563 34554 1.19 7.4E-01 BF34566.1 BST_HUMAN 21669 7.4E-01 BF346266.1 BST_HUMAN 21863 3.763 1.68 7.4E-01 AB021490.2 NT 21964 7.4E-01		13238	25712		7.5E-01		Ę	Homo saplens FRA3B common fragile region, diadenosine triphosphate hydrolase (FHIT) gene, excn 5
24318 5.26 7.5E-01 AF163151.2 NT 24638 30897 1.31 7.5E-01 D90907.1 NT 13771 24638 1.36 7.4E-01 AF162538.1 NT 16389 22854 0.63 7.4E-01 AF162538.1 NT 20328 33234 1.03 7.4E-01 AL168248.2 NT 20328 33235 1.03 7.4E-01 AL168248.2 NT 21107 34026 0.64 7.4E-01 AL16856.2 NT 21186 34492 7.17 7.4E-01 BF34626.1 EST_HUMAN 21863 34554 1.19 7.4E-01 BF34626.1 NT 22795 35786 0.59 7.4E-01 BF34626.1 NT 23964 37034 1.68 7.4E-01 AR164583.2 NT 24096 37660 7.4E-01 AR021490.2 NT 24096 37660 7.4E-01 AR021490.2 NT 24096 <td>↓_</td> <td>20050</td> <td>32923</td> <td></td> <td>7.5E-01</td> <td></td> <td>Ę</td> <td>Drosophila melanogaster tyrosine kinase receptor protein (eph) mRNA, complete cds</td>	↓_	20050	32923		7.5E-01		Ę	Drosophila melanogaster tyrosine kinase receptor protein (eph) mRNA, complete cds
24638 30897 1.91 7.5E-01 D90907.1 NT 13771 26279 1.36 7.4E-01 AI598146.1 EST_HUMAN 16389 28854 0.93 7.4E-01 AI185246.2 NT 20328 33234 1.03 7.4E-01 AL185246.2 NT 20328 33235 1.03 7.4E-01 AL181551.2 NT 21107 34026 0.69 7.4E-01 BF346266.1 EST_HUMAN 21166 34554 1.19 7.4E-01 BF346266.1 EST_HUMAN 21679 37034 1.68 7.4E-01 BR3492 NT 22795 35786 0.59 7.4E-01 BR3492 NT 23964 37034 1.68 7.4E-01 BR3490 NT 24096 7.4E-01 AB021490.2 NT NT 24096 7.4E-01 AB021490.2 NT NT 24096 7.4E-01 AB021490.2 NT NT 24096	!	24318		5.28			L	Homo sapiens dentin sialophosphoprotein precursor (DSPP) gene, complete cds
13771 26279 1.36 7.4E-01 AI598146.1 EST_HUMAN 16385 28854 0.95 7.4E-01 AF112538.1 NT 20328 33234 1.03 7.4E-01 AL161551.2 NT 20328 33235 1.03 7.4E-01 AL161551.2 NT 21107 34028 0.69 7.4E-01 BF346266.1 EST_HUMAN 21563 34492 7.17 7.4E-01 BF346266.1 EST_HUMAN 21563 34554 1.19 7.4E-01 BF346266.1 EST_HUMAN 1760.9 357034 1.68 7.4E-01 BB746266.1 EST_HUMAN 1760.9 357034 1.68 7.4E-01 BB746266.1 EST_HUMAN 1760.9 32043 37034 1.68 7.4E-01 AB021490.2 NT 24096 32043 37034 1.68 7.4E-01 AB021490.2 NT 24096 1.7804 2.93 7.4E-01 AB021490.2 NT 24096 1.7804 2.93 7.4E-01 AB021490.2 NT 24096 1.7804 2.93 7.7E-01 AB021490.2 NT 24096 1.7804 2.93 7.7E-01 AB021490.2 NT 24096 1.7804 2.93 7.7E-01 AB021490.1 NT 24096 1.7804 2.93 7.7E-01 AB021490.1 NT 25008 32942 7.77 7.3E-01 AD011418.1 NT 25008 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 E\$T_HUMAN 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 E\$T_HUMAN 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3E-01 AB0811.1 NT 250068 32943 7.77 7.3	 	24638					FZ	Synechocystis sp. PCC6803 complete genome, 9/27, 1056467-1188885
16389 28854 0.93 7.4E-01 AF112538.1 NT 16985 29430 7.7 7.4E-01 AL183246.2 NT 20328 33234 1.03 7.4E-01 AL181551.2 NT 20328 33234 1.03 7.4E-01 AL181551.2 NT 21107 34026 0.83 7.4E-01 BF346268.1 EST_HUMAN 21563 34492 7.17 7.4E-01 BF346268.1 EST_HUMAN 21563 34554 1.19 7.4E-01 BF346268.1 EST_HUMAN 22795 35786 0.59 7.4E-01 BF347503.1 EST_HUMAN 22795 35786 0.59 7.4E-01 AB021490.2 NT 23984 37034 1.68 7.4E-01 AB021490.2 NT 24096 7.4E-01 AB021490.2 NT NT 24176 25748 0.72 7.4E-01 AB021490.1 NT 17380 25839 2.93 7.3E-01 AB021490.1 NT 177823 30248 0.72 7.3E-01 AB021490.1 NT 17823	<u> </u>	13771	28279		7.4E-01	AI598146.1	EST HUMAN	h14b09.x1 NCI_CGAP_Bm25 Homo sepiens cDNA clone IMAGE:2187577 3' similar to contains Alu repetitive element; contains element MIR repetitive element
16985 29430 7.7 7.4E-01 AL163246.2 NT 20328 33234 1.03 7.4E-01 AL161551.2 NT 20328 33235 1.03 7.4E-01 BF346266.1 EST_HUMAN 21107 34026 0.69 7.4E-01 BF346266.1 EST_HUMAN 21563 34492 7.17 7.4E-01 BF346266.1 EST_HUMAN 21619 34554 1.19 7.4E-01 BF346266.1 EST_HUMAN 22795 35788 0.59 7.4E-01 BF346266.1 EST_HUMAN 23964 37034 1.68 7.4E-01 BE747503.1 EST_HUMAN 23964 37034 1.68 7.4E-01 AB021490.2 NT 24096 7.4E-01 AB021490.2 NT NT 24096 7.4E-01 AB021490.2 NT 17304 2.0748 0.72 7.4E-01 AB021490.2 NT 17304 2.0748 0.72 7.4E-01 AB021490.1 NT 17304 2.0748 0.72 7.2E-01 AE001480.1 NT 17823 3.0248	Ļ	16389	28854		7.4E-01		NT	Mava pusilla actin (Act1) mRNA, complete cds
20328 33234 1,03 7,4E-01 AL161551.2 NT 20328 33235 1,03 7,4E-01 AL161551.2 NT 21107 34026 0,83 7,4E-01 BF346266.1 EST_HUMAN 21186 0,64 7,4E-01 BF346266.1 EST_HUMAN 21563 34492 7,17 7,4E-01 BF747503.1 EST_HUMAN 22795 35788 0,59 7,4E-01 AL17263.1 EST_HUMAN 23964 37034 1,68 7,4E-01 AB021490.2 NT 24096 7,4E-01 AB021490.2 NT NT 24096 4,11 7,4E-01 AB021490.2 NT 24096 7,4E-01 AB021490.2 NT 17304 25748 0,72 7,4E-01 AL72641.1 NT 17304 25748 0,72 7,4E-01 AL72641.1 NT 17304 25748 0,72 7,3E-01 AL72641.1 NT 17823 30248 0,99 7,3E-01 AL72641.1 NT 18314 32117 5,86 7,3E-01 AL72641.1 NT	╄	16985	29430		7.4E-01	AL163246.2	NT	Hamo sapiens chromosome 21 segment HS21C046
20328 33235 1,03 7,4E-01 AL181551.2 NT 21107 34026 0,83 7,4E-01 BF346268.1 EST_HUMAN 21563 34492 7,17 7,4E-01 BF346268.1 EST_HUMAN 21563 34554 1,19 7,4E-01 BE747503.1 EST_HUMAN 22795 35786 0,59 7,4E-01 AA187988.1 EST_HUMAN 23964 37034 1,18 7,4E-01 AB021490.2 NT 24096 7,4E-01 AB021490.2 NT NT 24096 7,4E-01 AB021490.2 NT NT 24175 7,4E-01 AB021490.2 NT NT 24175 1,18 7,4E-01 AB021490.2 NT 24175 1,28 7,4E-01 AB021490.2 NT 24175 1,28 7,4E-01 AB021490.1 NT 17380 2,9839 2,93 7,3E-01 AE725421.1 NT 17823 30248 0,73 7,3E-01 AE725421.1 NT 18314 32117 5,86 7,3E-01 AE725421.1 NT	╙	20328	33234				FZ	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 51
21107 34028 0.63 7.4E-01 BF346266.1 EST_HUMAN 21186 34492 7.17 7.4E-01 U87990.1 NT 21563 34492 7.17 7.4E-01 BE747503.1 EST_HUMAN 22795 35784 1.19 7.4E-01 AA187986.1 EST_HUMAN 23964 37034 1.68 7.4E-01 AB021490.2 NT 23964 37035 1.68 7.4E-01 AB021490.2 NT 24036 37035 1.68 7.4E-01 AB021490.2 NT 24175 29748 0.72 7.4E-01 AB021490.2 NT 24175 29748 0.72 7.4E-01 AB021490.2 NT 17380 20839 2.93 7.3E-01 AA172641.1 NT 17823 30248 0.72 7.3E-01 AE01486.1 NT 17823 30248 0.89 7.3E-01 AB0141418.1 NT 20068 32942 7.77 7.3E-01	╙	20328			7.4E-01		L	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 51
21186 0.64 7.4E-01 U87990.1 NT 21563 34492 7.17 7.4E-01 BE747503.1 EST_HUMAN 21619 34554 1.19 7.4E-01 BE747503.1 EST_HUMAN 22795 35788 0.59 7.4E-01 AA187986.1 EST_HUMAN 23964 37034 1.68 7.4E-01 AB021490.2 NT 24096 37035 1.68 7.4E-01 AB021490.2 NT 24175 29748 0.72 7.4E-01 AB021490.2 NT 24175 29748 0.72 7.4E-01 AB021490.2 NT 17380 20839 2.93 7.3E-01 AA172641.1 NT 17823 20839 2.93 7.3E-01 AA50168.1 NT 17823 30248 0.99 7.3E-01 AA51725421.1 NT 18314 32117 5.86 7.3E-01 AA5172.1 NT 20068 32942 7.77 7.3E-01 AA51725421.1 <td>1_</td> <td>21107</td> <td></td> <td></td> <td></td> <td>BF346266.1</td> <td>EST_HUMAN</td> <td>602018456F1 NCI_CGAP_Brn67 Homo sapiens cDNA clone IMAGE:4154340 5'</td>	1_	21107				BF346266.1	EST_HUMAN	602018456F1 NCI_CGAP_Brn67 Homo sapiens cDNA clone IMAGE:4154340 5'
21563 34492 7.17 7.4E-01 BE747503.1 EST_HUMAN 22795 35786 0.59 7.4E-01 A1424933 NT 11424933 NT 23964 37034 1.68 7.4E-01 AB021490.2 NT 23964 37035 1.68 7.4E-01 AB021490.2 NT 24096 37035 1.68 7.4E-01 AB021490.2 NT 24096 4.11 7.4E-01 AB021490.2 NT 24096 4.11 7.4E-01 AB021490.2 NT 17304 29748 7.4E-01 AB021490.2 NT 17304 29748 7.4E-01 AB021490.2 NT 17304 29748 0.72 7.4E-01 AB021490.1 NT 17314 29748 0.72 7.3E-01 AF25421.1 NT 19314 32116 5.86 7.3E-01 AF25421.1 NT 19314 32117 5.86 7.3E-01 AD141418.1 NT 20068 32942 7.77 7.3E-01 AD141418.1 NT 20068 32943 7.77 7.3E-01 AM26511.1	L	21186		0.64	7.45-01		Ę	Rattus norvegicus leukocyte common antigen receptor (LAR) gene, trans-spliced alternative untranslated exon
21619 34554 1.19 7.4E-01 AA187986.1 EST_HUMAN 22795 35788 0.59 7.4E-01 11424833 NT 23964 37034 1.68 7.4E-01 AB021490.2 NT 23964 37035 1.68 7.4E-01 AB021490.2 NT 24096 4.11 7.4E-01 AB021490.2 NT 24076 4.11 7.4E-01 AB021490.2 NT 17304 29748 7.4E-01 AB021490.2 NT 17304 29748 7.3E-01 AB021490.2 NT 17304 29748 7.3E-01 AB021490.2 NT 17318 29839 2.93 7.3E-01 AF225421.1 NT 19314 32116 5.86 7.3E-01 A310372.1 NT 19314 3217 5.86 7.3E-01 A310141418.1 NT 20068 32942 7.77 7.3E-01 A30141418.1 NT 20068 32943 7.7	┺-	21563	34492		7.4E-01	BE747503.1	EST_HUMAN	801573028F1 NIH_MGC_9 Homo sapiens cDNA clone IMAGE:3834174 5'
22795 35786 0.59 7.4E-01 11424833 NT 23964 37034 1.68 7.4E-01 AB021480.2 NT 24096 4.11 7.4E-01 AB021480.2 NT 24175 1.28 7.4E-01 6783217 NT 24175 1.28 7.4E-01 AF22547 NT 17304 29748 0.72 7.3E-01 AF225421.1 NT 17823 30248 0.99 7.3E-01 AF225421.1 NT 18314 32116 5.86 7.3E-01 A3103 SWISSPROT 19314 32116 5.86 7.3E-01 A3103 SWISSPROT 19314 32117 5.86 7.3E-01 A3572.1 NT 20068 32942 7.77 7.3E-01 A2011418.1 NT 20068 32943 7.77 7.3E-01 AA878019.1 R\$T_HUMAN 23800 38859 3.86 7.3E-01 AA878019.1 E\$T_HUMAN	↓	21619			7.4E-01		EST HUMAN	487h01.s1 Stratagene endothelial cell 837223 Homo sapiens cDNA clone IMAGE:825297 3' similar to SW:TCPQ, MOUSE P42932 T-COMPLEX PROTEIN 1, THETA SUBUNIT
23964 37034 1 68 7 4E-01 AB021490.2 NT 23964 37035 1 68 7 4E-01 AB021490.2 NT 24096 4 11 7 4E-01 AB021490.2 NT 24175 4 11 7 4E-01 AB021490.2 NT 17304 29748 7 7E-01 AE001196.1 NT 17388 29639 7 3E-01 AE001196.1 NT 17823 30248 0 99 7 3E-01 AF225421.1 NT 18314 32116 5 86 7 3E-01 AF272.1 NT 18314 32116 5 86 7 3E-01 A5772.1 NT 20068 32942 7 77 7 3E-01 A2611.1 NT 20068 32942 7 77 7 3E-01 A26511.1 NT 20068 32943 7 77 7 3E-01 A26511.1 NT 23800 38659 3 36 7 3E-01 AA678019.1 EŞT_HUMAN	┺	22795			7.4E-01	11424933	NT	Homo sapiens NY-REN-45 antigen (LOC51133), mRNA
23984 37035 1.68 7.4E-01 AB021480.2 NT. 24096 4.11 7.4E-01 6753217 NT 24175 1.28 7.4E-01 A172841.1 EST_HUMAN 17304 29748 0.72 7.3E-01 AE001168.1 NT 17823 30248 0.72 7.3E-01 AF225421.1 NT 18314 32116 5.86 7.3E-01 L35772.1 NT 18314 32117 5.86 7.3E-01 L35772.1 NT 20068 32942 7.77 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 20068 32943 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 AAB78019.1 E\$T_HUMAN		23964	37034				N	Oryzias latipes gene for membrane guanyly cyclase OIGC1, complete cds
24096 4.11 7.4E-01 6753217 NT 24175 1.28 7.4E-01 AI472841.1 EST_HUMAN 17304 29748 0.72 7.3E-01 AE001168.1 NT 17823 30248 0.99 7.3E-01 AF225421.1 NT 19314 32116 5.86 7.3E-01 L35772.1 NT 18314 32117 5.86 7.3E-01 L35772.1 NT 20068 32942 7.77 7.3E-01 AD011418.1 NT 20068 32943 7.77 7.3E-01 AZ0511.1 NT 20068 32943 7.77 7.3E-01 AZ0511.1 NT 23800 38659 3.86 7.3E-01 AA078019.1 E\$T_HUMAN	⊢	23964	37035			AB021490.2	L	Oryclas latipes gene for membrane guanylyl cyclase OIGC1, complete cds
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17304 29748 0.72 7.3E-01 AE001168.1 NT 17388 29839 2.93 7.3E-01 AF225421.1 NT 17823 30248 0.99 7.3E-01 O43103 SWISSPROT 19314 32116 5.86 7.3E-01 L35772.1 NT 24777 32525 0.82 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 20068 32943 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 AA978019.1 E\$T_HUMAN	ι.	24175		1.28	7.4E-01	AI472641.1	EST_HUMAN	ta13h01.x1 NCI_CGAP_Lym5 Homo sapiens cDNA clone IMAGE:2043985 3'
17388 29839 2.93 7.3E-01 AF225421.1 NT 17823 30248 0.99 7.3E-01 O43103 SWISSPROT 19314 32116 5.86 7.3E-01 L3572.1 NT 24777 32525 0.82 7.3E-01 A5772.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 20068 32943 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 A878019.1 E\$T_HUMAN		17304			7.3E-01	AE001166.1	NT	Borrelia burgdorferi (section 52 of 70) of the complete genome
17823 30248 0.99 7.3E-01 O43103 SWISSPROT 19314 32116 5.86 7.3E-01 L35772.1 NT 19314 32117 5.86 7.3E-01 L35772.1 NT 24777 32525 0.82 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 AAB78019.1 E\$T_HUMAN	L.	17388			7.3E-01	AF225421.1	N	Homo sapiens HT017 mRNA, complete cds
18314 32116 5.86 7.3E-01 L35772.1 NT 18314 32117 5.86 7.3E-01 L35772.1 NT 24777 32525 0.82 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 AAB78019.1 E\$T_HUMAN		17823					SWISSPROT	FERRICHROME SIDEROPHORE PEPTIDE SYNTHETASE
19314 32117 5.86 7.3E-01 L35772.1 NT 24777 32525 0.82 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M28511.1 NT 20068 32943 7.77 7.3E-01 M28511.1 NT 23800 38859 3.86 7.3E-01 AA878019.1 EŞT_HUMAN	<u> </u>	19314	32116			L35772.1	N	Mus musculus antigen (CD72) gene
24777 32525 0.82 7.3E-01 AJ011418.1 NT 20068 32942 7.77 7.3E-01 M26511.1 NT 20068 32943 7.77 7.3E-01 M26511.1 NT 23800 38859 3.86 7.3E-01 AA678019.1 EŞT_HUMAN	-	19314					IN	Mus musculus antigen (CD72) gene
20068 32842 7.77 7.3E-01 M26511.1 NT 20068 32943 7.77 7.3E-01 M26511.1 NT 23800 38859 3.86 7.3E-01 AA678019.1 EŞT_HUMAN	L	24777	32525				IN	Lycopersicon esculentum mRNA for ubiquitin ectivating enzyme
20068 32943 7.77 7.3E-01 M26511.1 NT 23800 38859 3.86 7.3E-01 AA678019.1 EŞT_HUMAN	با	20068			7.3E-01		IN	V. alginolyticus sucrase (scrB) gene, complete cds
23800 36859 3.86 7.3E-01/AA678019.1 [EŞT_HUMAN	느	20068			7.3E-01	M26511.1	NT	V. alginolyticus sucrase (scrB) gene, complete cds
	L	23800			7.3E-01	AA678019.1	EŞT_HUMAN	2/25b08.s1 Scares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA clone IMAGE:431799 3'

Page 38 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Page 39 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Expression Signal(Top) Hit AvalueTop Hit Acassion No.Top Hit Descriptor SourceTop Hit Descriptor Source	Į.	1.3 7.0E-01 AB014514.1 INT Homo sapiens mRNA for KIAA0614 protein, partial cds	1.22 7.0E-01/N62412.1 EST HUMAN contains Alu repetitive element.	1.22 7.0E-01/N62412.1 EST_HUMAN contains Alu repetitive element.	1,98 7,0E-01 AL163301.2 NT Homo sapiens chromosome 21 segment HS21C101	2.99 7.0E-01 AE003921.1 INT Xylelia fastidiosa, section 67 of 229 of the complete genome	1.03 7.0E-01 AB021316.1 NT Arabidopsis thallana mRNA for chlorophyl b synthase, complete cds	11.92 7.0E-01 AE000253.1 NT Escherichia odi K-12 MG1655 section 143 of 400 of the complete genome		0.81 7.0E-01 U53868.1 NT and mtID genes, complete cds		7.0E-01 U53868.1 NT	1.99 7.0E-01 AV763842.1 EST_HUMAN AV763842 MDS Homo sapiens cDNA cione MDSCHE04 5'	1.99 7.0E-01 AV763842.1 EST_HUMAN AV763842 MDS Homo sepiens cDNA clone MDSCHE04 5'	7.0E-01 9630464	10.2 6.9E-01 U69874.1 NT cds	Candida albicans squalene epoxidase (CAERG1) gene, complete cds and translational regulator gene, partial	저	T_HUMAN	1.7 6.9E-01/AE002271.2 NT Chlamydle mundarum, section 3 of 85 of the complete genome	0.8 6.9E-01/AB035692.1 NT Brenchiostoma betcheri BbNA3 mRNA for notochord ectin, complete cds	1.3.1 6.9E-01 BE298188.1 [EST_HUMAN 601177333F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:3532328 5'	3.4 6.9E-01 AL 161573.2 NT Arabidopsis thallana DNA chromosome 4, contig fragment No. 69	3.4 6.9E-01/AL161573.2 NT Arabidopsis thaliana DNA chromosome 4, contig fragment No. 69	0.83 6.9E-01 AF118046.1 NT Entamoeba disper cation transporting ATP ase (atpase) gene, partial cds		0.62 6.9E-01 AF206319.1 NT Musa acuminata pectate lyase 1 (PL1) mRNA, complete cds	EST_HUMAN	8.9E-01 D89013.1 NT	1.94 6.9E-01 D89013.1 NT Homo sapiens DAN gene, complete cds
Most Similar (Top) Hit BLASTE Value									,	7.0E-01								6.9E-0	0-36-0	0-36'9										6.9E-(
Expression Signal	1.3	1.3	1.22	1.22	1.98	2.99	1.03	11.92		0.61		0.61	1.99	1.99	1.35	10.2		10.2	2.8	1.7	0.8	1.31	3.4	3.4	0.83	0.62	0.62	0.68	1.94	1.94
ORF SEQ ID NO:		26388	27630	27631		30336				34714			36546	36547		26130			28472		31310	31893				35073	35074	L		38888
Exon SEQ ID NO:	13868		15057	15057	17778	17922	18723	20860		21768	•	21788	23513	23513		13616		13616	13948	15868	18576	19108	20463	20463	21634	22111		1		23646
Probe SEO ID NO:	1272	1272	2492	2492	5213	5362	8107	8319		9240		9240	10999	10999	12594	1005		1005	1353	3256	5954	8208	7921	7921	8606	1	1196	10307	11138	11138

Page 40 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Mus muscutus major histocompatibility complex region NG27, NG28, RPS28, NADH oxidoreductase, NG29, KiFC1, Fas-binding protein, BING1, tapasin, RalCDS-like, KE2, BING4, beta 1,3-galactosy transferase, and Mus musculus major histocompatibility complex region NG27, NG28, RPS28, NADH oxidoreductase, NG29, KIFC1, Fas-binding protein, BING1, tapasin, RalGDS-like, KE2, BING4, beta 1,3-galactusyl transferase, and Homo sapiens nuclear factor of kappa light polypeptide gene enhancer in B-cells 1 (NFKB1) gene, complete Homo sapiens nuclear factor of kappa light potypapitde gene enhancer in B-cells 1 (NFKB1) gene, complete ORKHEAD BOX PROTEIN C2 (FORKHEAD-RELATED PROTEIN FXHL14) (MESENCHYME FORK aj75e05.s1 Soares_parathyroid_tumor_NbHPA Homo sapiens cDNA clone IMAGE:1402256 3' similar to nv13e07.s1 NCI_CGAP_Pr22 Homo sapiens cDNA clone IMAGE:1220100 3' similar to gb:X13546_ma1 Drosophila melanogaster Mst85C gene, complete cds; NMDMC isoform (Nmdmc) gene, complete cds, Quail fast skeletal muscle troporin I gene, complete cds EXT2g12.s1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone IMAGE.786310 3' similar to contains element TAR1 repetitive element; Anopheles gambiae strain M2 translation initiation factor 4C (1A) (eIF-4C) mRNA, complete cds alternatively spliced; and transcription factor (Relish) gene, complete cds, alternatively spliced gb:X58411_ma1 ALCOHOL DEHYDROGENASE CLASS II PI CHAIN (HUMAN); HEAD PROTEIN 1) (MFH-1 PROTEIN) (TRANSCRIPTION FACTOR FKH-14) wn31f02.x1 NCI_CGAP_Gas4 Homo sapiens cDNA clone IMAGE:2447087 3 Synechocystis sp. PCC6803 complete genome, 27/27, 3418852-3573470 Human HMG-17 gene for non-histone chromosomal protein (HUMAN) Mus musculus Wiskott-Aldrich syndrome protein (Wasp), mRNA Stagonospora avenae bgl1 gene for beta-glucosidase, exons 1-4 Stagonospora avenae bgl1 gene for beta-glucosidase, exons 1-4 Top Hit Descriptor Mus musculus zinc finger protein (Peg3) mRNA, complete cds Mus musculus zinc finger protein (Peg3) mRNA, complete cds S.tuberosum mRNA for glucose-6-phosphate dehydrogenase Giardia intestinalis carbamate kinase gene, complete cds Homo sapiens mRNA for KIAA1345 protein, partial cds RPS18 genes, complete cds; Sacm21 gene, partial> Rat(hooded) protectin gene: exon iii and flanks RPS18 genes, complete cds; Sacm21 gene, ĝ ş EST_HUMAN EST HUMAN EST_HUMAN EST_HUMAN SWISSPROT Database 世合 Source ¥ Ż 눋 Ż 눋 Ħ Ł Ę Ę 눋 달토토 ż 6678580 Top Hit Acession 6.7E-01 AF213884.1 6.7E-01 AF186073.1 6.7E-01 AA451864.1 6.8E-01 AF110520.1 6.8E-01 AF110520.1 AAB54475.1 6.8E-01 AA687936.1 AJ276675.1 6.8E-01 AF038939.1 AF038939.1 AF213884. 6.9E-01 AI888312.1 ģ 6.8E-01 AB037766. AF164151 6.8E-01 D90917.1 6.7E-01 M12132. 6.7E-01 X74421. 6.8E-01 J00762.1 8.9E-01 Q99958 6.8E-01 6.8E-01 6.8E-01 6.7E-01 6.7E-01 6.8E-01 6.BE-01 (Top) Hit BLAST E Vost Similar 4.28 2.18 0.64 1.77 27.63 26.51 18 0.48 2.96 1.77 2.36 1.33 1.28 1.62 1.45 2.96 Expression Signal 28120 29581 36505 36506 36540 25493 27340 27361 36995 36996 25463 ORF SEQ ID NO: 26783 29706 30611 26118 36541 35017 13010 15460 15642 17133 23480 23925 23925 14768 22756 12974 14539 24870 14249 23480 23507 25003 13604 15255 22056 23507 SEO ID ġ 3026 4550 10965 10965 11475 11475 361 2192 2211 2856 10993 320 SEQ ID 12670 2698 4672 9556 10261 10993 11651 992

Page 41 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 42 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

	Exan NC: NC: NC: 17784 17897 19415 22733 23050 23102 23102 23102 23102 23102 23103 23103 23103 23103 23103 23103 23103 23103 1187 1774 1778 1778 1778 1778 1778 1778	ORF SEQ ID NO: 30202 30312 32231 33062 35261 28583 28619 28689 28619 28689 28619 28619 28619 28619 28619 35501 35501 35501 35501 35501 35501 35501 35501 35501	Signa Signa	Most Similar (Top) Hit BLAST E Value 6.5E-01 6		Top Hit Deftabase Source Source Source Source Source EST HUMAN EST HUMAN INT INT INT INT INT INT INT INT INT IN	Pheseolus vulgaris ATPase gamma subunit mRNA, nuclear gene encoding mitochondrial protein, pertral cots H septers mRNA for inmunoglobulin heavy chain variable region (904-46 VH4, 4-59/DP-71) Chicken mRNA for inmunoglobulin heavy chain variable region (904-46 VH4, 4-59/DP-71) Chicken mRNA for inmunoglobulin heavy chain variable region (904-46 VH4, 4-59/DP-71) Chicken mRNA for IntS-Dar meterosomal matrix protein, complete ods Wd5802x1 NOIC_CGAP_P-P/2B Homo saplers cDNA chore INAGE: 100748 3' W47106.1 Soares placenta_8toSweeks_2NDH-P8109W Homo saplers cDNA done IMAGE: 100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 1100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 1100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 1100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 1100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 1100748 3' AU138078 PLACE I Hono saplers cDNA chore INAGE: 179130 3' Scerevisse chronosome IV reading frame ORF YDL097c Drosophila malanogastes Rd dynain light chain mRNA, complete cds Mmusculus dystroglycan I UPACI, grane, exorn 1 and 2 and complete cds Hono saplers above learning cetta and septems cDNA chore INAGE: 4291126 5' HISTIDINE-RICH PROTEIN PRECURSOR (CLONE PPHRP-III) Hearnophilis influenzae Rd socion 4' of 18 of complete genome Treponena pallidum section 83 of 37 of the complete genome Theorems pallidum section 83 of 37 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 83 of 57 of the complete genome Theorems pallidum section 84 of 57 of 50 o
3050 6214 6712	111	1 {	0.7			EST_HUMAN NT	Lycopes score rescurentum poses gene, comprete COS PMO-BT0757-0105500-202-405 BT0757 Homo sapiens cDNA Streptococcus dysgelecties (meg) gene, complete ads Streptococcus dyscretarities (meg) nene complete ads
6712	19306	32111	_	6.3E-01	27788.1	<u>z</u> .	Streptococcus dysgalactiae (mag) gene, comptete cds

PCT/US01/00669

Page 43 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

WO 01/57277

Top Hit Descriptor	601676889F1 NIH_MGC_21 Hama sapiens cDNA clane IMAGE:3959351 5'	glycoprotein IIIa (Alu 1 and 3 fusion junction) [human, Genomic Mutant, 300 nt]	601884050F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4102596 5'	Varida virus, complete genome	Varida virus, complete genome	Chlamydia muridarum, section 59 of 85 of the complete genome	S.cerevisiae chromosome VII reading frame ORF YGR218w	Escherichia coli K-12 MG1655 section 203 of 400 of the complete genome	nr09h06.s1 NCI_CGAP_Co10 Hamo sapiens cDNA done IMAGE:11613713' similar to TR:002916 002916	HLARK.;	CM-BT043-090289-046 BT043 Hamo sapiens cDNA	HYPOTHETICAL 13.7 KD PROTEIN IN INO1-IDS2 INTERGENIC REGION	HYPOTHETICAL 15.3 KD PROTEIN IN VMA12-APN1 INTERGENIC REGION	Mus musculus keratin complex 2, gene 8g (Krt2-8g), mRNA	Homo sapiens 3'-phosphoadenosine 5'-phosphosulfate synthetase (PAPSS) mRNA, complete cds	C.limicola pscD gene	Spermophilus susilicus isolate S47 cytochrome b (cytb) gene, complete cds; mitochondrial gene for	mitochondrial product	HYPOTHETICAL 142.5 KD PROTEIN C23E2.02 IN CHROMOSOME I	Mus musculus calcium-sensing receptor related protein 4 (Casr-rs4) mRNA, partial cds	Mus musculus chromosome X contigA; putative Magea9 gene, Celtractin, NAD(P) steroid dehydrogenase	and zinc ringer process 165	ysureversity some spiedal living spiedal living sepieds curve curve my one to be and debut some many	Lycycla skyni esturation by the outer to superconductors generally and the complete of superconductors generally and superconductors are superconductors.	601336146F1 NIH_MGC_44 Homo sapiens cDNA clone IMAGE:3690010 5	Human pulmonary surfactant-associated protein SP-B (SFTP3) mRNA, complete cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 23	NON-STRUCTURAL POLYPROTEIN [CONTAINS: RNA-DIRECTED RNA POLYMERASE; THIOL	PROTEASE P3C; HELICASE (2CLIKE PROTEIN); COAT PROTEIN]	NON-STRUCTURAL POLYPROTEIN (CONTAINS: RNA-DIRECTED RNA POLYMERASE ; THIOL PROTEASE P3C : HELICASE (2011 KE PROTEIN); COAT PROTEIN)	Ne misculus secretar acidic extens rich obconstella (Sparc) mRNA	the Higherman sections strain of the Basel control (percent of the Basel control of the Base	Homo sapiens soute carrier family 20 (surate transporter), member 2 (SEC 2042) mixing
Top Hit Databese Source	EST_HUMAN 6	NT B	T_HUMAN			NT C	NT	NT.	<u>-</u>			SWISSPROT	SWISSPROT		¥	Į	8		SWISSPROT I	TN TN			EST_HUMAN	<u> </u>	T HUMAN				SWISSPROT	TOGGSSIMS			
Top Hit Acession No.	BE902044.1	S62927.1	BF216984.1	9627521 NT	9627521 NT	AE002329.2	273003.1	AE000313.1		AA877715.1	AI904160.1	P47003	P36073	9910293 NT	AF105227.1	X83528.1		AF157898.1	Q10135	AF022253.1		2	H72255.1	AF034411 1	Ī		2		P27410	022440	TIA SCOOTSS	0/09/00	4557538 NT
Most Similar (Top) Hit BLAST E Value	6.3E-01		6.3E-01	6.3E-01	6.3E-01		6.3E-01			6.3E-01		6.3E-01	6.3E-01	6.3E-01						6.2E-01			6.2E-01		6.2E-01	6.2E-01	6.2E-01		6.2E-01	P 20 04	0.45.04	8.TE-U1	6.1E-01
Expression Signal	3.32	16:0	1.15	2.9	2.9	79.0	1.52	0.87		2.45	15.21	1.94	2.02	30.63	1.85	3.2		0.71	2.03	3.14		1.08	5.65	27.0	1.75	2.35	5.85		3.76	ar c	3.70	4.95	1.05
ORF SEQ ID NO:	T	34284	34627	34804	34805			35915		36479		36853						30171	31390	l			33703		33212		35472		35927		07850		29666
Exon SEQ 1D NO:	86602	21358	21682	21855	21855	22349	22818	22915		23456		L	23908	25042	<u> </u>		1	17742	ı	20028	l	- 1	20784		20309		1_	L_	22923	1	1	- 1	17215
Probe SEQ ID NO:	8458	8819	9147	9341	9341	1986	10324	10421		10939	11216	11302	11458	11769	11864	12082		5175	6030	7506		7548	8243	0020	9370	9429	0666		10429	3,6,	10469	2438	4632

Page 44 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 45 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
10174	22689		1.61	6.0E-01	Q01497	SWISSPROT	PEROXISOMAL MEMBRANE PROTEIN PER9 (PEROXIN-3)
10936	23453	36476	2.14		AJ131892.1	NT	Gallus gallus mRNA for Hyperion protein, 419 kD isoform
10936	23453	36477	2.14	8.0E-01	AJ131892.1	1N	Gallus galtus mRNA for Hyperion protein, 419 kD isoform
11426	23877	36942	2.84	6.0E-01	AI420623.1	EST_HUMAN	#08/07.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:2095621 3'
12158	24398					LNT	Homo saplens nuclear factor (erythroid-derived 2)-like 3 (NFE2L3), mRNA
12265	24475		1.99		AA70608	EST_HUMAN	296g05.s1 Soares_fetal_liver_spleen_1NFLS_S1 Homo saplens cDNA clone IMAGE:462776 3
12426	24879		1.29	6.0E-01	5803136 NT	LN	Homo sapiens RNA binding motif protein 3 (RBM3), mRNA
12469	24885	30709		6.0E-01		L	Mus musculus cGMP-inhibited phosphodiesterase (Pde3a), mRNA
12499	24810		6.92		BE157617.1	EST_HUMAN	RC1-HT0375-030500-015-c03 HT0375 Homo saplens cDNA
1038	13648	26160	1.09	5.9E-01	U32701.1	L	Haemophilus influenzae Rd section 16 of 163 of the complete genome
1447	14039	26568	1.06			NT	Mus musculus 3-hydroxy-3-methylgiutaryl-Coenzyme A lyase (Hmgcl), mRNA
3308	15919	28395	5.12	5.95-01	AL163267.2	LΝ	Homo sapiens chromosome 21 segment HS21C067
3308	15919	28396	5.12		AL163267.2	LN	Homo sapiens chromosome 21 segment HS21C067
4304				5.9E-01	AF162756.1	NT	Rattus norvegicus cenexin 2 mRNA, partial cds
							also deliberate boson a como (COCIII) II e la describa se la como deliberate boson deliberate la como deliberate boson delibe
659	- !	1				L	Homo capiens low density lipoprotein receptor-related protein II (LRFZ) gene, exch 1 and partual cus
7310	19838					L	Homo sepiens gene for histamine HZ receptor, promoter region and complete cds
7941	20483	33395	0.57	5.9E-01	D90911.1	Ä	Synechacystis sp. PCC6803 camplets genome, 13/27, 1576593-1719643
9462	21887	34943	0 83	5.96-01	AF063204.2	¥	Chlamydia trachomatis strain K/UW31/Cx major outer membrane protein (omp1) gene, complete cds
9827					-	SWISSPROT	E6 PROTEIN
10091	上	35579	1.15	5.9E-01	P55284	SWISSPROT	VASCULAR ENDOTHELIAL-CADHERIN PRECURSOR (VE-CADHERIN) (CADHERIN-5)
10551	上		3.24		Q9X0I3	SWISSPROT	THYMIDYLATE KINASE (DTMP KINASE)
10557	23083	36105		5.9E-01	AF197944.1	TN	Xenopus laevis receptor protein tyrosine phosphatase delta (XPTP-D) mRNA, complete cds
10840	23361		3	L	AW837175.1	EST_HUMAN	PM1-DT0041-190100-002-h03 DT0041 Homo sapiens cDNA
11073	23585	36626	2.25		AF064626.1	NT	Mus spretus strain SPRET/Ei CD48 antigen (Cd48) gene, partial cds
11810	24182	31030	1.92	5.9E-01	L42320.1	LΝ	Oryctologus cuniculus alpha 1 anti-trypsin (alpha 1 AT) gene, promoter region
12053	24336		2.88	5.9E-01	AB017705.1	TN	Aspergillus cryzae pyrG gene for orotidine-5'-phosphate decarboxylase, complete cds
12280	24483		7.56		P34926	SWISSPROT	MICROTUBULE-ASSOCIATED PROTEIN 1A [CONTAINS: MAP1 LIGHT CHAIN LC2]
1952	14536	27092		5.8E-01	P40472	SWISSPROT	SIM1 PROTEIN
4056	16653	29119	1.22	5.8E-01		EST_HUMAN	601852474F1 NIH_MGC_56 Hamo seplens CDNA clone IMAGE:4076131 5
4612	17195	29641	3.73			FZ	Vigna radiata mRNA for proton pyrophosphatase, complete cds
4914			1.18	١		LZ.	Megaselia scalaris sex-lethal homolog (Megsxl) gene, partial cds, atternatively spliced products
5577	18208		0.75	5.8E-01	AE002152.1	N-	Ureaplasma urealyticum section 53 of 59 of the complete genome

Page 46 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	Single Excit 10555 Expressed III otal EIVO	Most Similar Top Hit Acession (Top) Hit Acession ORF SEQ Expression (Top) Hit Descriptor Top Hit Descriptor Signal No. Signal Value Source	31051 2.52 5.8E-01 Q10699 SWISSPROT	31713 2.37 5.8E-01 D78659.1 EST_HUMAN	31840 0.71	9837 2.47 5.8E-01 S65091.1 NT cyclic AMP-regulated phosphoprotein (rats, mRNA, 1030 nt)	0370 2.57 5.8E-01 H41571.1 EST HUMAN gb:S78187 M-PHASE INDUCER PHOSPHATASE 2 (HUMAN):	33477 0.66 5.8E-01 AI280051.1 EST HUMAN	33478 0.66 5.8E-01 AI280051.1 EST_HUMAN	33582 2.34 5.8E-01 P14328 SWISSPROT	33583 2.34 5.8E-01 P14328 SWISSPROT	34287 9.48 5.8E-01/AJ270774.1 NT	34363 0.88 5.8E-01 Q27368 SWISSPROT	34364 0.56 5.8E-01 020471 SWISSPROT	0.89 5.8E-01 BF031606.1 [EST_HUMAN	36405 9.44 5.8E-01 AJ243213.1 NT	3.66 5.8E-01[BF700092.1 [EST_HUMAN	2.04 5.8E-01 BF700092.1 [EST_HUMAN	0.66 5.7E-01 6755253 NT	28352 1.58	2.63 5.7E-01 AB033503.1 NT	6571 29041 3.09 5.7E-01 AF011581.1 NT Homo sepiens T cell receptor beta chain (BV6S7*2-BJ1S1) mRNA, partial cds	31881 3.67 5.7E-01 BF035413.1 EST_HUMAN	32219 0.72 5.7E-01 AA194201.1 EST_HUMAN	30478 1.28 5.7E-01 AL111440.1 NT	P00373 SWISSPROT	TN	22213 35186 1.17 5.7E-01 AL161532.2 NT Arabidopsis thaliana DNA chromosome 4. config fragment No. 32	35187 1.17 5.7E-01	35973 0.86 5.7E-01 BF540962.1	28498 1 5.6E-01 AB018283.2 NT	28499 1 5.6E-01 AB018283.2 NT	16910 29351 0.69 5.6E-01 D83135,1 NT Chicken TBP gene, exon8, complete cds
															11			2				L	L		L		3						
I ACTU INIMIMIAL EIMIMIMIMIMIMIMIMIMIMIMIMI MIMIMIMIMIMI			722 18348	331 18937	454 19055	903 19637			031 20573	1131 20672	1131 20672		902 21440	903 21441	1514 22014	869 23390	915 23434	021 23535	15694	15872	1552 16158	1973 16571	19097	3812 19403	3945 18053	L	L	L	1			L	4324 1691

Page 47 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	SEQ ID	ORF SEQ ID NO:	Expression	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Datebase Source	Top Hit Descriptor
8738	21277	34201	4.42		AV684703.1	EST_HUMAN	AV684703 GKC Hamo sapiens cDNA clane GKCFSF05 5'
9297			1.11	5.6E-01	AB038782.1	IN	Homo sapiens MUC3A gene for intestinal mucin, partal cds
11658			2.5	5.6E-01	BE888280.1	EST_HUMAN	601514007F1 NIH_MGC_71 Homo sapiens cDNA clone IMAGE:3915457 5'
							ng75g10.s1 NCI_CGAP_Pr6 Homo sepiens cDNA clone IMAGE.940674 similar to centains element PTR7
11779	24168	36775	1.28	5.6E-01	AA493535.1	EST_HUMAN	repetitive element;
12156		30490	3.31		1.2	NT	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 13
12185	1_		2.56	5.6E-01	P50505	SWISSPROT	HIGH AFFINITY POTASSIUM TRANSPORTER
12619	24698		3.11	5.6E-01	BF573829.1	EST_HUMAN	602132029F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4271334 5
1253	l.,	26367	1.13	5.5E-01	8393912 NT	TN	Rattus norvegicus Propionyl Coenzyme A carboxylase, beta polypeptide (Pccb), mRNA
27.25		27847	13.6	5.5E-01	P03341	SWISSPROT	GAG POLYPROTEIN (CONTAINS: INNER COAT PROTEIN P12: CORE PROTEIN P15; CORE SHELL PROTEIN P30; NUCLEOPROTEIN P10]
	L.						GAG POLYPROTEIN (CONTAINS: INNER COAT PROTEIN P12; CORE PROTEIN P15; CORE SHELL
2725		}		5.5E-01	P03341	SWISSPROT	PROTEIN P30; NUCLEOPROTEIN P10]
2943		28033	0.69	5.5E-01	2902085 NT	- 1	Homo sapiens superkiller viralicidic activity 2 (S. cerevisiee nomolog)-like (SKIVZL), mKNA
3102	15717		1.51	5.5E-0.	H46219.1	EST_HUMAN	yo18a10.s1 Soares adult brain N2b5HB55Y Homo saplens cDNA clone IMAGE:178268 3
3271	15883	28365	2.68	-0-35'S	AF227240.1	LN	Rabbit oral papillomavirus, complete genome
3755	16356		0.97	5.5E-0	P48755	SWISSPROT	FOS-RELATED ANTIGEN-1
8388	L.,	33846	0.66	5.5E-01	A1791,788.1	EST_HUMAN	or82c01.y5 NCI_CGAP_Lu5 Homo sapiens cDNA clone IMAGE:1802338 5
9682			0.74	5.5E-01	U88415.1	INT	Crimean-Congo hemorrhagic fever virus strain SPU 415/85 nucleoprotein gene, complete cds
10279	<u> </u>	35763	0.84	5.5E-0	T05047.1	EST_HUMAN	EST02935 Fetal brain, Strategene (catt936206) Homo sapiens cDNA clone HFBCQ35
151	12814	25301	12.97	5.4E-01	7657266 NT	NT	Homo saplens KIAA0929 protein Msx2 Interacting nuclear target (MINT) homolog (KIAA0929), mRNA
151	12814	25302	12.97	5.4E-01	7657266 NT	N	Homo sapiens KIAA0929 protein Msx2 interacting nuclear target (MINT) homolog (KIAA0929), mRNA
3	<u> </u>			_	6 4E 01 A E232008 1	Ę	Pseudomonas syringae pv. tornato strain DC3000 AvrE (avrE). HrpW (hrpW), and GstA (gstA) genes, complete cds: and unknown genes
01.5	13239	61/67	2.		AI ESECUCI.		Barrell And Get (Get (and Get (ask) and Get (as
611	13239	25714	1.6		5.4E-01 AF232008.1	NT	r seudomorais syringae pr. toriado suain Dobodo Anie, aniejo, impro, anie con tarro, general complete edis, and unknown genes
1314			~		5.4E-01 AW896087.1	EST_HUMAN	QV4-NN0040-070400-160-c04 NN0040 Hamo sapiens cDNA
2154	ı				5.4E-01 AE002247.2	TN	Chlamydophila pneumoniae AR39, section 74 of 94 of the complete genome
2286	14870	27446	2.18		AJ276682.1	NT	Drosophila melanogaster mRNA for 15,15' beta carotene dioxygenase (beta-diox gene)
3984	16582	29053	0.62		5.4E-01 U07561.1	NT	Human ABL gene, exon 1b and intron 1b, and putative M8804 Met protein (M8604 Met) gene, complete cds
5259					5.4E-01 AW747972.1	EST_HUMAN	QV0-BT0041-061099-033-602 BT0041 Homo sapiens cDNA

Page 48 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Vatue	Top Hit Acession No.	Top Hit Detabese Source	Top Hit Descriptor
5388	13239	25713	0.59	5.4E-01	AF232006.1	LN	Pseudomonas syringae pv. tomato strain DC3000 AvrE (avrE), HrpW (hrpW), and GstA (gstA) genes, complete cds; and unknown genes
5388	13239	25714	0.59	5.4E-01	AF232008.1	LΝ	Pseudomonas syringae pv. tomato strain DC3000 AvrE (avrE), HrpW (hrpW), and GstA (gstA) genes, complete cds; and unknown genes
5838	18462	L			AW842327.1	T_HUMAN	PM2-CN0030-030200-003-c10 CN0030 Homo sapiens cDNA
6338	18944	31723	1.49	5.4E-01	AB025017.1	NT	Rattus norvegicus gene for 71511, complete cds
7094	19665	32504	1.1		BE966592.2	EST_HUMAN	601660276R1 NIH_MGC_71 Hamo sapiens cDNA clone IMAGE:3906090 3'
7374	19900		0.75		221619.1		S.cerevisiae RIB3 gene encoding DBP synthase
7374	19900	32763	0.75	5.4E-01	221619.1	LN	S. cerevisiae RIB3 gene encoding DBP synthase
							MITOCHONDRIAL TRIFUNCTIONAL ENZYME ALPHA SUBUNIT PRECURSOR (TP-ALPHA) [INCLUDES: LONG-CHAIN ENOYL-COA HYDRATASE ; LONG CHAIN 3-HYDROXYACYL-COA
7376	19902	32768	1.47	5.4E-01	064428	SWISSPROT	DEHYDROGENASE]
980	22398		1.98	5.4E-01	BF572536.1	EST_HUMAN	602076545F1 NIH_MGC_62 Homo sapiens cDNA clone IMAGE:4243890 5'
10957	23472	36497	3.25	5.4E-01	P36858	SWISSPROT	NITRATE REDUCTASE [NADPH] (NR)
11485	23934	37004		5.4E-01	Q60675	SWISSPROT	LAMININ ALPHA-2 CHAIN PRECURSOR (LAMININ M CHAIN) (MEROSIN HEAVY CHAIN)
11485	l	37005	5.79	5.4E-01	Q60675	SWISSPROT	LAMININ ALPHA-2 CHAIN PRECURSOR (LAMININ M CHAIN) (MEROSIN HEAVY CHAIN)
11586	18944	31723	2.42	5.4E-01	AB025017.1	NT	Rattus norvegicus gene for TIS11, complete cds
11725	24132		2.52	5.4E-01	AI858398.1	EST_HUMAN	wi37g04.x1 NCI_CGAP_Ut1 Homo sapiens cDNA clone IMAGE:2427126 3' similar to gb:M13452 LAMIN A (HUMAN);
	l						Homo sapiens HLA class III region containing tenascin X (tenascin-X) gene, partial cds; cytochrome P450 21-
542	13173	25653	2.29	5.3E-01	AF019413.1	Ę	(Bf), and complement component C2 (C2) genes.>
2811	15363	27931		5.3E-01		F	Homo sapiens protein tyrosine phosphatase, receptor-type, zeta polypeptide 1 (PTPRZ1) mRNA
2811	15363	27932	6.51	5.3E-01	4506328 NT	LN TN	Homo sapiens protein tyrosine phosphatase, receptor-type, zeta polypeptide 1 (PTPRZ1) mRNA
3280	15891	L	3.13			NT	Homo sapiens secreted C-type lectin precursor (LSLCL) gene, complete cds
4290	16876		1.39	5.35-01	U39687.1	NT	Mycoplasma genitalium section 9 of 51 of the complete genome
5649	18277	30753	1.91	5.3E-01	A1820921.1	EST_HUMAN	zu42h12.y5 Soares ovary tumor NbHOT Hamo sapiens cDNA clone IMAGE:740711 5'
5649	18277	L	1.91	5.3E-01	AI820921.1	EST_HUMAN	2u42h12.y5 Soares ovary tumor NbHOT Homo sapiens cDNA clone IMAGE:740711 5'
5742	18368	31075	0.87	5.3E-01	AA193672.1	EST_HUMAN	z 42g09.r1 Soares_NhHMPu_S1 Homo sapiens cDNA clone IMAGE:668112 5
5742	18368	31076	0.87	5.3E-01	AA193672.1	EST_HUMAN	zr42g09.r1 Soares_NhHMPu_S1 Homo sapiens cDNA clone IMAGE:666112.5
5827	18451	31174	1.84	5.3E-01	BE645620.1	EST_HUMAN	7e73c12.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:3288118 3' similar to gb:302783 PROTEIN DISULFIDE ISOMERASE PRECURSOR (HUMAN);
5827	18451	31175	1.84	5.3E-01	BE645620.1	EST HUMAN	7e73c12.x1 NCI_CGAP_P/28 Homo septens cDNA clone IMAGE:3288118 3' similar to gb:J02783 PROTEIN DISULFIDE ISOMERASE PRECURSOR (HUMAN);
	١			J			

Page 49 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
8835	21374		1.83	5.3E-01	L01950.2	±N	Roridula gorgonias ribulose 1,5-bisphosphate carboxylase (rbcL) gene, partial ods; chloroplast gene for chloroplast product
8885	21423	34348	8.0	5.3E-01	BF433956.1	EST_HUMAN	7q71c12.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE: 3' similar to contains element MER29 repetitive element;
8885	21423	34349	9.0	5.3E-01	BF433956.1	EST_HUMAN	7q71c12.x1 NCI_CGAP_Lu24 Homo sepiens cDNA clone IMAGE: 3' similer to contains element MER29 repetitive element;
10112	22607	35597	0.48	5.3E-01	A1954210.1	EST_HUMAN	wx84b02.x1 NCI_CGAP_Mel15 Homo sapiens cDNA clone IMAGE:2551275 3' similar to SW:COXA_HUMAN P20874 CYTOCHROME C OXIDASE POLYPEPTIDE VA PRECURSOR;
11435	23885		6.92		BE566291.1	EST_HUMAN	601339867F1 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3882188 5'
11650	24881		4.22	5.3E-01	AA916053.1	EST_HUMAN	og30e05.s1 NCI_CGAP_Br7 Homo sapiens cDNA clone IMAGE:1441378 3' similar to gb:J02611 APOLIPOPROTEIN D PRECURSOR (HUMAN);
849	13465	25973	19.16	5.2E-01	1.20770.1	NT	Drosophila melanogaster helix-loop-helix mRNA, complete cds
1206	1	26319	10.07		Q9WV30	SWISSPROT	NUCLEAR FACTOR OF ACTIVATED T CELLS 5 (T CELL TRANSCRIPTION FACTOR NFAT5) (NF-AT5) (REL DOMAIN-CONTAINING TRANSCRIPTION FACTOR NFAT5)
1233	13832	26346	2.91	5.2E-01	AF224492.1	LN	Homo sapiens phospholipid scramblese 1 gene, complete cds
1930	14514		4.11	5.2E-01	AL163285.2	NT	Homo sapiens chromosome 21 segment HS21C085
2191			2.97		AB018283.2	NT	Homo sapiens mRNA for KIAA0740 protein, partial cds
3153	15767	28233	1.67	5.2E-01	U65942.1	NT	Chlamydophila abortus strain S26/3 POMP91A and POMP90A precursor, genes, complete ods
3274	15886		0.71		D73443.1	INT	Azotobacter vinelandii icd gene for isocitrate dehydrogenase, complete cds
3452			1.74	5.2E-01		TN	Botrytis cinerea strain T4 cDNA library under conditions of nitrogen deprivation
3492	16097	28572	2.49		AA984165.1	EST_HUMAN	am77g05.s1 Stratagene schizo brain S11 Homo sapiens cDNA clone IMAGE:1616504 3'
7 091	18205	•	20 0	5 2F-01	AF020269 1	5	Medicago sativa chloroplast malate dehydrogenase precursor (p1mdh) mRNA, nuclear gene encoding chloroplast protein, complete cds
5161			0.87			¥	Mus musculus vanilloid receptor-like protein 1 (Vrt1), mRNA
5314	17876		66'0	5.2E-01	AL163281.2	Z	Homo sapiens chromosome 21 segment HS21C081
5834	18458	31179	76.0		AA284261.1	EST_HUMAN	2c44d09.T7 Soares, senescent_fibroblasts_NbHSF Homo sapiens cDNA clone IMAGE:325169 3'
9846	24795	35115	1.19	L	X02218.1	Ę	Chicken duplicated genes for histone H2A, H4 and a histone H3 gene
9846	24795	35116	1.19	5.2E-01	X02218.1	Z	Chicken duplicated genes for histone H2A, H4 and a histone H3 gene
9845	22343	35325	0.64		AA194518.1	EST_HUMAN	2q05b09.r1 Stratagene muscle 937209 Homo sapiens cDNA clone IMAGE:628793 5
9940	22435	35411	1.65	5.2E-01	AF143952.2	NT	Homo sapiens PELOTA (PELOTA) gene, complete cds
12590	24682		4.94	5.2E-01	P18516	SWISSPROT	RETINOIC ACID RECEPTOR GAMMA (RAR-GAMMA) (RETINOIC ACID RECEPTOR DELTA) (RAR- DELTA)
645	13268	25746	2.13		M58509.1	Z-	Human adrenodoxin reductase gene, exons 3 to 12
878	H	1		Ц	AJ233944.1	NT	Polyangium vitellinum (strain Pl w1) 16S rRNA gene

Page 50 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	Τορ Hit Descriptα	Pdyangium vitellinum (strain PI vt1) 16S rRNA gene	R.norvegicus mRNA for mammalian fusca protein	602139319F1 NIH_MGC_46 Homo saplens cDNA clone IMAGE:4298117 5	w/39b12.x1 NCI_CGAP_Ut1 Homo sapiens cDNA clone IMAGE:24272633'	TRANSCRIPTION-REPAIR COUPLING FACTOR (TRCF)	IL2-BT0731-250400-077-G08 BT0731 Homo sapiens cDNA	AV712328 DCA Homo sapiens cDNA clone DCAAUF07 5'	y94a09.s1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:146872 3'	QV4-ST0023-160400-172-a01 ST0023 Homo sapiens cDNA	QV4-ST0023-160400-172-e01 ST0023 Homo sapiens cDNA	Human regenerating protein (reg) gene, complete cds	65B1 Human retina cDNA Tsp509I-cleaved sublibrary Homo saplens cDNA not directional	Human carboxyl ester lipase (CEL) gene, complete cds	801556863F1 NIH_MGC_58 Homo sapiens cDNA clone IMAGE:3828767 5'	nac51f10.x1 NCI_CGAP_Bm23 Homo saplens cDNA clone IMAGE:3406218 3' similar to contains element	TAR1 repetitive element;	Homo sapiens postmeiotic segregation increased 2-like 9 (PMS2L9), mRNA	Homo sapiens postmeiotic segregation increased 2-like 9 (PMS2L9), mRNA	Buchnera aphidicola genomic fragment containing (chaperone Hsp80) groEL, DNA biosynthesis initiating protein (dnsA), ATP operon (atpCDGAHFEB), and putative chromosome replication protein (gidA) genes,	complete cds; and termination factor Rho (rho) gene>	Buchnera aphidicola genomic fragment containing (chaperone Hsp&0) groEL, DNA biosynthesis initiating protein (dnAA). ATP coeron (atbCDGAHFEB), and putative chromosome replication protein (gidA) genes.	complete cds, and termination factor Rho (rho) gene>	Thermotoga maritima section 97 of 138 of the complete genome	Mus musculus anti-DNA immunoglobulin light chain IgM mRNA, antibody 363p.138, partial cds	Homo saplens mRNA for KIAA1184 protein, partial cds	Xenopus laevis smooth muscle beta-tropomyosin mRNA, complete cds	601823850R1 NIH_MGC_79 Horno sapiens cDNA clone IMAGE:40434853'	601903871F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:4136632 5	GLYCOGEN DEBRANCHING ENZYME (GLYCOGEN DEBRANCHER) [INCLUDES: 4-ALPHA-GLUCANOTRANSFERASE); AMYLO-1,8-GLUCOSIDASE	(DEXTRIN 6-ALPHA-D-GLUCOSIDASE))
Personal Lines Explanation	Top Hit Database Source	Į.	FN	EST_HUMAN	EST_HUMAN	SWISSPROT	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	⊢ Z	EST_HUMAN		EST_HUMAN		EST_HUMAN	LN	NT		NT		Z.	NT	IN	LN.	NT	EST_HUMAN	EST_HUMAN		SWISSPROT
oligio.	Top Hit Acession No.	AJ233944.1	X87885.1	BF683095.1	AI858495.1	P96380	BE091796.1	AV712326.1	R80873.1	AW806881.1	AW 806881.1	J05412.1		5.1E-01 M94579.1	1 BF030207.1		1 BF 439982.1	488552 NT	488552 NT		5.0E-01 AF008210.1		5.0E-01 AF008210.1	5.0E-01 AE001785.1	5.0E-01 U55574.1	5.0E-01 AB033010.1	M92304.1	5.0E-01 BF107848.1	5.0E-01 BF317212.1		P35573
	Most Similar (Top) Hit BLAST E Value	5.1E-01	5.1E-01	5.1E-01		5.1E-01	5.1E-01	5.1E-01	5.1E-01	5.1E-01	5.15-01	5.1E-01	5.1E-01	5.1E-01	5.1E-01		5.1E-01	5.0E-01	5.0E-01		5.0E-01		5.0E-01	5.0E-01	5.0E-01	5.0E-01			5.0E-01		5.0E-01 P35573
	Expression Signal	3.98	0.88	11.33	4.61	3.03	17.0	0.79	1.42	0.73	0.73	4.6	3.4	0.95	2.04		2.01	1.4	1.4		5.48		5.48	5.58	0.65	3.11	1.78	0.64	3.1		1.34
	ORF SEQ ID NO:	25782				28300			32316	33968		35065						27326	27327		27335		27336		28875					_	35004
	Exon SEQ ID NO:	13300	14284	l	16743	16852	17793	19025	19495	21048		L		22560	L	<u> </u>	24385	14757	14757		14785		14785	16341	16410	16540		Į.	l	L	22043
	Probe SEQ ID NO:	876	1692	2069	4151	4266	5229	6422	6997	8507	8507	9602	9605	10065	11874		12129	2180	2180		2189		2189	3740	3811	3942	8487	8604	9379		9543

Page 51 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	GLYCOGEN DEBRANCHING ENZYME (GLYCOGEN DEBRANCHER) (INCLUDES: 4-ALPHA-GLUCANOTRANSFERASE), AMYLO-1,8-GLUCOSIDASE (DEXTRIN 8-ALPHA-D-GLUCOSIDASE))	601445024F1 NIH_MGC_65 Homo sapiens cDNA clone IMAGE:3849438 5	Mus musculus MRC OX-2 antigen homolog gene, exons 2-5, and complete cds	Homo sapiens chromosome 21 segment HS21C102	NUCLEAR ENVELOPE PROTEIN CUT11	602076648F1 NIH_MGC_62 Homo sapiens cDNA clone IMAGE:4243860 5'	Xenopus laevis mRNA for c-Jun protein, 1978 BP	Cavia porcellus pulmonary surfactant protein A (SP-a) mRNA, complete cds	FIBRILLIN 1 PRECURSOR	Homo sapiens discylglycerol kinase 3 (DAGK3) gene, exon 10	Homo sapiens diacy/glycerol kinase 3 (DAGK3) gene, exon 10	Oryza sativa subsp. japonica mEF-G mRNA for mitochondrial elongation factor G, complete cds	601874964F1 NIH_MGC_54 Homo sapiens cDNA clone IMAGE:4102503 5'	hc90c02.x1 Soares_NFL_T_GBC_S1 Homo sapiens*cDNA clone IMAGE:2907268 3' similar'to TR:099714 095714 HERC2.;	Mus musculus unc13 homolog (C. elegans) 1 (Unc13h1), mRNA	Mus musculus adenylyl cyclase 1 (Adcy1) cDNA, partial cds	Homo sapiens neurotrophin-1/B-cell stimulating factor-3 gene, complete cds	nq22e11.s1 NCI_CGAP_Co10 Homo sapiens cDNA clone IMAGE:1144652 3'	Homo sapiens chromosome 21 segment HS21C101	Homo sapiens eukaryotic translation initiation factor 4 gamma, 1 (EIF4G1), mRNA	ol32a09.s1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:1525144 3'	Homo sapiens potassium channel, subfamily K, member 5 (TASK-2) (KCNK5) mRNA, and translated	products	Saccharomycas cerevislae) sporulation protein (SPO11) gene required for meiotic recombination, complete	cds	nu85f09.s1 NCI_CGAP_Alv1 Homo sapiens cDNA clone IMAGE:1217513	Homo sapiens reproduction 8 (D8S2298E) mRNA	Hamo sapiens chramasame 21 segment HS21C009	Arabidopsis thallana DNA chromosome 4, contig fregment No. 4	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 4
Top Hit Database Source	SWISSPROT	EST_HUMAN	NT	TN	SWISSPROT	EST_HUMAN	NT	IN	SWISSPROT	LΝ	۲Z	ĽN	EST_HUMAN	EST HUMAN	Z.	F	Ę	EST_HUMAN	卢	TN	EST_HUMAN		N T		M	EST_HUMAN	N	NT	TN	Z
Top Hit Acession No.	P35573	BE869218.1	AF029215.1	AL163302.2	013961	BF571462.1	AJ243955.1	U40869.1	Q61554	AF020931.1	AF020931.1	AB040051.1	BF209791.1	AW339905.1	10946863 NT	AF053980.1	AF176912.1	AA613562.1	AL163301.2	11431438 NT	AA912842.1		4504850 NT		J02987.1	AA659878.1	5031650 NT	AL163209.2	AL161492.2	AL161492.2
Most Similar (Top) Hit BLAST E Value	5.0E-01	5.0E-01	5.0E-01	5.05-01	5.0E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	_	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.9E-01	4.8E-01		4.8E-01		4.8E-01	4.8E-01	4.8E-01	4.8E-01		4.8E-01
Expression Signal	1.34	1.04	3.45	2.38	4.27	2.31	1.6	1.35	1.32	2.35	2.35	1.9	1.49	96.0	2.2	0.74	2.48	5.73	1.74	1.36	1.05		0.62	1	8.6	4.22	1.85	0.87	3.72	3.72
ORF SEQ ID NO:	35005					25946	26827	27089	30681	31565	31568	32862		34592		35706			30872						30827			33061	33138	
Exan SEQ ID NO:	22043	22786	24187	24656	24668	13439	14292	14533	18231	18797	18797	19997	21458	21651		22715	24117	25081	24657	24708	16195		17011		18324	18381	19883	20174	20246	1
Probe SEQ ID NO:	9543	10291	11815	12554	12569	822	1699	1949	2095	6187	6187	7475	8920	9115	9220	10220	11704	12546	12555	12630	3591		4782		5698	6790	7357	7662	7738	7738

Page 52 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

		MAGE:154795 5' similar to contains element		E:4300048 5'					E:4096387 5	4GE:1755544 3	811 5 end	811 5'end		ne IMAGE:2581580 3'	ta cds	16, and partial cds	MAGE:4181303 5		E:3912488 5	one IMAGE:2909198 3	nt. position (7/7)		5E:4245481 5	SE:4245481 5	spo ex		UNIT (EPITHELIAL NA+ CHANNEL GAMMA JM CHANNEL 1 GAMMA SUBUNIT)		3E:4129472 5	3E:4129472 5'		12 10000	3E:3843637 5
Single Exult Flubes Expressed in Feditary	Top Hit Descriptor	y77f10.y5 Soares breast 2NbHBst Homo sapiens cDNA clone IMAGE:154795 5' similar to contains element MER6 repetitive element ;	PM1-HT0350-201299-004-b04 HT0350 Homo sapiens cDNA	602184267F1 NIH_MGC_42 Homo sapiens cDNA clone IMAGE:4300048 5	S.cerevisiae ORFs from chromosome X	Homo sapiens chromosome 21 segment HS21C027	Trypanosoma cruzi transposon VIP II SIRE repeat region	Chlamydomonas reinhardtii cop gene, exons 1-8	601883880F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4096387 5	qf72a09.x1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:1755544 3	hbc811 Human pancreatic islet Homo sapiens cDNA clone hbc811 5'end	hbc811 Human pancreatic islet Homo sapiens cDNA clone hbc811 5'end	Rattus norvegicus Spermine binding protein (Sbp), mRNA	xb69e11.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2581580 3	Influenza A virus isolate hk51697 hemagglutinin (HA) gene, partial cds	Human collegen alpha2(XI) (COL11A2) gene, exons 6 through 16, and partial cds	602043889F1 NCI_CGAP_Bm67 Homo sapiens cDNA clone IMAGE:4181303 5	RC6-NT0029-240400-011-E08 NT0029 Homo sepiens cDNA	601511333F1 NIH_MGC_71 Homo sapiens cDNA clone IMAGE:3912488 5	hd11c08 x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2909198 3	Pyrococcus horikoshii OT3 genomic DNA, 1485001-1738505 nt. position (717)	RC1-ST0278-040400-018-b06 ST0278 Homo sapiens cDNA	602081103F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:42454815	602081103F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4245481 5	Bovine steroid 21-hydroxylase gene (P-450-c21) gene, complete cds	Homo saptens chromosome 21 segment HS21C048	AMILORIDE-SENSITIVE SODIUM CHANNEL GAMMA-SUBUNIT (EPITHELIAL NA+ CHANNEL GAMMA SUBUNIT) (GAMMA ENAC) (NONVOLTAGE-GATED SODIUM CHANNEL 1 GAMMA SUBUNIT)	(SCNEG) (GAMMA NACH)	601900234F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:4129472 5	601900234F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:4129472 5	INTERFERON REGULATORY FACTOR 3 (IRF-3)	INTERFERON REGULATORY FACTOR 3 (IRF-3)	601568755F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3843637 5
EXOII FIODES	Top Hit Database Source	EST_HUMAN	EST_HUMAN	EST_HUMAN	NT	TN	LN	ΝΤ	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	ĮN	EST_HUMAN	NT	IN	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	ΙN	EST_HUMAN	EST_HUMAN	EST_HUMAN	INT	LN		SWISSPROT	EST_HUMAN	EST_HUMAN	SWISSPROT	SWISSPROT	EST_HUMAN
alfaile	Top Hit Acession No.	1 A1820744.1	4.8E-01 BE155148.1	1 BF568633.1	X83502.1	4.8E-01 AL163227.2	4.8E-01 AF227565.1	4.8E-01 AJ132984.1	4.7E-01 BF217173.1	4.7E-01 AI204374.1	4.7E-01 T11414.1	4.7E-01 T11414.1	6981501 NT	AW087791.1	4.7E-01 AF102673.1	4.7E-01 U41069.1	4.7E-01 BF529658.1	4.7E-01 AW889448.1	1 BE887763.1	1 AW341561.1	11 AP000007.1	4.6E-01 AW818638.1	4.6E-01 BF693300.1	4.6E-01 BF693300.1	1 M11267.1	31 AL163248.2		01 P51170	01 BF313593.1	4.6E-01 BF313593.1	01 Q90643	01 Q90643	01 BE734781.1
	Most Similar (Top) Hit BLAST E Value	4.8E-01	4.8E-01	4.8E-01	4.8E-01 X83502.1	4.8E-01	4.8E-01	4.8E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.7E-01	4.6E-01	4.6E-01	4.6E-01	4.6E-01	4.6E-01		4.6E-01	4.6E-01	4.6E-01	4.6E-01	4.6E-01	4.6E-01
	Expression Signal	1.36	1.13	0.58	2.02	1.29	3.04	3.36	8.72	0.78	0.52	0.52	0.5	0.79	4.94	2.19	11.61	2.89	1.92	1.33	1.38	2.23	1.68	1.68	1.03	22.08		1,37	1.12	1.12	3.27		2.39
	ORF SEQ ID NO:	33291		-					32036	32263	33257	l		35972		36503	36715					28862	28870	28871		30390		30400		L			31067
	SEQ ID NO:	20388	21748	22417	23141	24170	24842	24895	19234	19447	20349			22961	l		L		24243	ı	1_			16406	L	17986		17995	ı			i i	18361
	Probe SEQ ID NO:	7846	9169	9921	10807	11786	12016	12846	9839	7107	7806	7806	9008	10467	10727	10963	11163	11254	11904	12036	12666	3797	3806	3806	5323	5429		5440	5612	5612	2863	5863	5735

Page 53 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

	ľ						
Probe SEQ ID NO:	Exen SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	. Top Hit Descriptor
5748	18374	31082	4.22	4.6E-01	AI247679.1	EST_HUMAN	qh59h02.x1 Soares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA clone IMAGE:18490113' similar to TR:015338 015338 BUTYROPHILIN ;
5748	18374	31083	4.22	4.6E-01	AI247679.1	EST_HUMAN	dh59h02.x1 Soares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA clone IMAGE:1849011 3' similar to TR:015338 015338 BUTYROPHILIN.
5756	18382	31094	1.4	4.6E-01	P20050	SWISSPROT	MEIOSIS SPECIFIC PROTEIN HOP1
5828			1.05	4.6E-01	AF212124.1	ΝŢ	Anolls schwartz cytochrome b gene, partial cds; mitochondrial gene for mitochondrial product
5907			98'0	4.6E-01	BE817247.1	EST_HUMAN	PM0-BN0260-120600-001-F07 BN0260 Homo sapiens cDNA
8 8 8 8	18675	31417	0.75	4.6E-01	D26215.1	NT	Unidentified soil bacteria 16S rRNA gene encoding 18S ribosomal RNA
6404	19007	31788	1.05	4,6E-01	AE000894.1	<u>_</u>	Methanobacterium thermosulotrophicum from bases 1165751 to 1176238 (section 100 of 148) of the complete cenome
6865	19589	32429	1.36	4.6E-01	U62332.1	Į.	Emericella nidulans NEMPA (nempA) gene, mitochondrial gene encoding putative mitochondrial protein, complete cds.
6865	19599	32430	1.36	4.6E-01	U62332.1	ŢN	Emericella nidulans NEMPA (nempA) gene, mitochondrial gene encoding putative mitochondrial protein, complete ods
7712		33108	98'0	4.6E-01	AA483577.1	EST HUMAN	nh04h05.s1 NCI_CGAP_Thyl Homo saptens cDNA clone IMAGE:943363 similar to contains Alu repetitive element.contains element 1 repetitive element:
8262	20803	33721	13.23	4.6E-01	BF697399.1	EST HUMAN	602130953F1 NIH_MGC_56 Homo sapiens cDNA clone IMAGE:4287828 5
9225	21741	34684	1.04	4.6E-01	P55202	SWISSPROT	ATRIAL NATRIURETIC PEPTIDE RECEPTOR B PRECURSOR (ANP-B) (ANPRB) (GC-B) (GUANYLATE CYCLASE)
9225	21741	34685	1.04	4.6E-01	P55202	SWISSPROT	ATRIAL NATRIURETIC PEPTIDE RECEPTOR B PRECURSOR (ANP-B) (ANPRB) (GC-B) (GUANYLATE CYCLASE)
9583	22083	35046	0.55	4.6E-01	AF162283.1	NT	Glycine max acetyl-CoA carboxylase (accB-1) gene, complete cds; nuclear gene for chloroplast product
9583	22083	35047	0.55	4.6E-01	AF162283.1	NT	Glycine max ecetyl-CoA carboxylase (accB-1) gene, complete cds; nuclear gene for chiproplast product
9886	22383	35358	2.63	4.6E-01	AI915634.1	EST HUMAN	wg73e12.x1 Scares NSF F8 9W OT PA P S1 Homo sapiens cDNA clone IMAGE:2370766 3
9886		35359	2.63	4.6E-01	Al915634.1	EST_HUMAN	wg73e12.x1 Scares NSF F8 9W OT PA P S1 Homo sapiens cDNA clone IMAGE:2370766 3
10870	23391		3.09	4.6E-01	P98163	SWISSPROT	PUTATIVE VITELLOGENIN RECEPTOR PRECURSOR (YL)
10879		36416	4.13	4.6E-01	BE185449.1	EST_HUMAN	L5-HT0730-100500-075-g05 HT0730 Homo sapiens cDNA
10879		38417	4.13	4.6E-01	BE185449.1	EST_HUMAN	L5-HT0730-100500-075-g05 HT0730 Homo sapiens cDNA
11348	23044	36054	5.52	4.6E-01	AF019369.1	NT	Human thiopurine methyttransferase (TPMT) gene, exon 10 and complete cds
11348	23044	36055	5.52	4.6E-01	AF019369.1	NT	Human thicpurine methyltransferase (TPMT) gene, exon 10 and complete cds
12654	24726		1.26	4.6E-01	M22360.1	TN	Ret plasma proteinase inhibitor alpha-1-inhibitor III group 3 variants 6J, 12J, 13J, and 17J mRNA, partial ods

Page 54 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO: 1954 1954 1954 2895 3346 3346 4160 4160 4160 4160 6719 6719 6719 7604	Exen SEQ ID NO: 14538 14538 15512 15	ORF SEQ ID NO: 27084 27085 27085 27085 28432 28434 28432 28432 28432 28432 30092 31070 31070	Signal 1.69 1.69 4.77 4.77 4.04 4.04 4.04 1.35 1.35 1.35 1.35 1.35 1.35 1.35 1.35	Most Similar (Top) Hit BLAST E Vatue 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01 4.5E-01	AW 608814 SE BESS 445. A RESS 48. A RESS 849. 1	Top Hit Detabase Source NT TOP HIT NT NT NT EST_HUMAN EST_HUMAN EST_HUMAN EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN NT EST_HUMAN EST_HUMAN EST_HUMAN NT EST_HUMAN EST_HUMAN EST_HUMAN NT EST_HUMAN EST_HU	Top Hit Detroccus radiodurars R1 section 68 of 229 of the complete chromosome 1 I EST HUMAN Application of the complete chromosome 1 EST HUMAN D'NAMIN-1 (HUMAN): Swissprot PRECURSOR (HSPG) (PERLECAN) (PLC) Swissprot POLAGEN APPA 5(IV) CHAIN SWISSPROT COLLAGEN APPA 5(IV) CHAIN EST HUMAN Resolute DNA polymerase epsilon catalytic subunit (Pole) gene, exons 2 though 12 EST HUMAN Resolute DNA polymerase epsilon catalytic subunit (Pole) gene, exons 2 though 12 EST HUMAN Resolute DNA polymerase epsilon catalytic subunit (Pole) gene, exons 2 though 12 EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits SWISSPROT COLLAGEN ALPHA 5(IV) CHAIN EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute DNA polymerase epsilon catalytic subunit gene, complete cits EST HUMAN Resolute protessome regulator PA28 beta subunit gene, complete cits WASSERROT Rat nucleolar proteins B23.1 and B23.2 WASSERROT Rat nucleolar proteins B23.1 and B23.2 WASSEROT Resolute Resolute Resolution subiens cDNA clone IMAGE:2426618 3' similar to TR:092023 09202
8342	1 1	33804	4.02		M32661.1 Al648596.1	NT EST_HUMAN	D.melanogaster Shaw2 protein mRNA, complete cds tz58g11.x1 NCI_CGAP_Ov35 Homo sapiens cDNA ctone IMAGE:2292644 3'
8494	21033	33954	0.69		Q52728	ISSPROT	POLY-BETA-HYDROXYBUTYRATE POLYMERASE (POLY(3-HYDROXYBUTYRATE) POLYMERASE) (PHB POLYMERASE) (PHB SYNTHASE) (POLY(3-HYDROXYALKANOATE) POLYMERASE) (PHA POLYMERASE) (PHA SYNTHASE) (POLYHYDROXYALKANOIC ACID SYNTHASE)
8929	21255	34385	1.74	_	11444786 NT		Homo sapiens hypothetical protein DKFZp547G183 (DKFZp547G183), mRNA
8853	22351	200	1.02	4.5E-01	AE000218.1 NT		Escherichia coli K-12 MG1655 section 108 of 400 of the complete genome
10389	11	35877	23.95	7	M86006.1	T HUMAN	EST02531 Fetal brain. Stratagene (rattg382/6) Homo caniene child closs UEBCV17
10389	22883	35878	23.95	4.5E-01	M86008.1	П	EST02531 Fetal brain, Stratagene (cattro30206) Homo sapiens CDNA clone HFBC 717
10744	23268	36285	3.01		_	EST_HUMAN	xo14h01.x1 NCI_CGAP_Ut3 Home sapiens cDNA clone IMAGE:2703985 3' similar to SW:INT6_MOUSE
11131	23639	1	1.9	4.5E-01	AV719382.1	EST_HUMAN	AV719382 GLC Hamo sapiens cDNA clone GLCCED12 5

WO 01/57277 PCT/US01/00669

Page 55 of 526 Table 4 Single Exon Probes Expressed in Feta

Fig. 2002 Control Co					ľ	aligie	EXON Propes	Single Exon Probes Expressed in Fetal Liver
NO: UNO: Signal BLAST EST HUMAN	Probe SEQ ID	Exon SEQ ID	ORF SEQ	Expression	Moet Similar (Top) Hit	Top Hit Acession	Top Hit Database	. Ton Hit Descriptor
23828 36890 1.68 4.5E-01 BE094451.1 EST HUMAN 24540 2.13 4.5E-01 BE071461.1 EST HUMAN 24540 2.13 4.5E-01 BF337531.1 EST HUMAN 24578 6.25 4.5E-01 F737531.1 EST HUMAN 14682 2.7572 3.26 4.4E-01 P69765 SWISSPROT 15965 28442 1.27 4.4E-01 AFD8790.1 NT 15965 28442 1.27 4.4E-01 AFD8790.1 NT 15965 28443 1.27 4.4E-01 BF141396.1 NT 15965 28446 2.31 4.4E-01 BF141396.1 RST HUMAN 17874 30693 4.06 4.4E-01 BF141396.1 EST HUMAN 18242 30693 4.06 4.4E-01 BF141396.1 RST HUMAN 18724 31476 1.53 4.4E-01 BF141396.1 RST HUMAN 18724 31476 1.53 4.4E-01 BF141396.1	Ö	ö	<u> </u>	Brigio	MAST E	o Ž	Source	and more than the second secon
25670 3.3 4.5E-01 BE871461.1 EST HUMAN 24540 2.13 4.5E-01 BF337531.1 EST HUMAN 24578 6.25 4.5E-01 BF337531.1 EST HUMAN 14989 27572 3.26 4.4E-01 BF058790.1 NT 15985 28442 1.27 4.4E-01 AF058790.1 NT 15986 28446 2.31 4.4E-01 BF058790.1 NT 15989 27572 3.26 4.4E-01 BF058790.1 NT 15989 28446 2.31 4.4E-01 BF058790.1 NT 15989 28446 2.31 4.4E-01 BF058790.1 NT 16804 1.28 4.4E-01 BF058790.1 NT 17706 2.07 4.4E-01 BE378707.1 EST HUMAN 18242 30693 4.06 4.4E-01 BE378707.1 EST HUMAN 18489 31215 1.72 4.4E-01 BE378707.1 EST HUMAN 18724 31476 1.53 4.4E-01 AV080399 SWISSPROT 18724 31477 1.53 4.4E-01 AV080399	11378			1.68	4.5E-01		EST HUMAN	RC3-BT0333-160300-016-e03 BT0333 Homo sapiens cDNA
24540 2.13 4.5E-01 BF33753.1 EST_HUMAN 2457B 6.25 4.5E-01 11422099 NT 11422099 NT 14989 27572 3.26 4.4E-01 B6805033 NT 15965 28443 1.27 4.4E-01 B7658790.1 NT 15965 28443 1.27 4.4E-01 B7658790.1 NT 15969 28446 2.31 4.4E-01 B6786726.1 EST_HUMAN 16804 1.27 4.4E-01 B6786726.1 EST_HUMAN 17804 2.07 4.4E-01 B6141396.1 EST_HUMAN 17804 2.07 4.4E-01 B6141396.1 EST_HUMAN 18242 30693 4.06 4.4E-01 B6141396.1 EST_HUMAN 18242 30693 4.06 4.4E-01 B625019.1 NT 18242 30693 4.06 4.4E-01 B6720408.1 EST_HUMAN 1874 1.53 4.4E-01 B7720408.1 EST_HUMAN 1874 31477	11671	25070		3.3	4.5E-01		EST_HUMAN	601449201F1 NIH_MGC_65 Homo sapiens cDNA clone IMAGE:3852961 5'
24578 6.25 4.5E-01 11422099 NT 14682 1.39 4.4E-01 6680503 NT 14682 27572 3.26 4.4E-01 6680503 NT 15965 28442 1.27 4.4E-01 AF058790.1 NT 15965 28443 1.27 4.4E-01 BF058720.1 EST HUMAN 15966 28443 2.31 4.4E-01 BF058720.1 EST HUMAN 1706 2.07 4.4E-01 BE143396.1 EST HUMAN 17706 2.07 4.4E-01 BE143396.1 EST HUMAN 17839 30265 0.94 4.4E-01 BE143396.1 EST HUMAN 18242 30692 4.06 4.4E-01 BF0829 SWISSPROT 18242 30693 4.06 4.4E-01 BF0829 SWISSPROT 18724 31476 1.53 4.4E-01 BF0829 SWISSPROT 18850 31771 1.69 4.4E-01 BF0829 SWISSPROT <	12370			2.13	4.5E-01		EST_HUMAN	602035275F1 NCI_CGAP_Bm84 Homo sapiens cDNA clone IMAGE:4183280 5'
14862 1.39 4.4E-01 6680503 NT 14869 27572 3.26 4.4E-01 P49765 SWISSPROT 15865 28442 1.27 4.4E-01 AF08760.1 NT 15865 28446 1.27 4.4E-01 BF086726.1 EST_HUMAN 16904 1.27 4.4E-01 BF086726.1 EST_HUMAN 1796 2.07 4.4E-01 BE141396.1 EST_HUMAN 17974 0.9 4.4E-01 BC141396.1 EST_HUMAN 18242 30692 4.06 4.4E-01 BAW14885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 BAW14885.1 EST_HUMAN 18724 31216 1.72 4.4E-01 BAW14885.1 EST_HUMAN 18724 31476 1.53 4.4E-01 BAW18885.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AW184885.1 EST_HUMAN 18805 31215 1.53 4.4E-01 AW194913.1 EST_HUMAN	12442			6.25	4.5E-01		N.	Homo sapiens testis-specific kinase 2 (TESK2), mRNA
14999 27572 3.26 4.4E-01 P49765 SWISSPROT 15965 28442 1.27 4.4E-01 AF058790.1 NT 15965 28443 1.27 4.4E-01 BF058726.1 EST HUMAN 15969 28446 2.31 4.4E-01 BF058726.1 EST HUMAN 1706 1.28 4.4E-01 BE141398.1 EST HUMAN 1706 2.07 4.4E-01 BE141398.1 EST HUMAN 17974 0.94 4.4E-01 BE141398.1 EST HUMAN 18242 30692 4.06 4.4E-01 BO4929 SWISSPROT 18242 30693 4.06 4.4E-01 BO4929 SWISSPROT 18724 31275 1.72 4.4E-01 BO4929 SWISSPROT 18724 31477 1.53 4.4E-01 AV720408.1 EST HUMAN 18724 31477 1.53 4.4E-01 AV720408.1 EST HUMAN 18963 32818 0.89 4.4E-01 AV720408.1	2081	14662		1.39	4.4E-01		Z	Mus musculus integral membrane-associated protein 1 (Itmap1), mRNA
15965 28442 1.27 4.4E-01 AF058790.1 NT 15965 28443 1.27 4.4E-01 AF058790.1 NT 15969 28446 2.31 4.4E-01 BF058728.1 EST_HUMAN 1706 2.07 4.4E-01 BF058728.1 EST_HUMAN 17839 30265 0.94 4.4E-01 BE141398.1 EST_HUMAN 1784 4.09 4.4E-01 BE141398.1 EST_HUMAN 18242 30692 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW614885.1 EST_HUMAN 18505 31231 1.53 4.4E-01 AV720408.1 EST_HUMAN 18506 31231 1.53 4.4E-01 AW680785.1 EST_HUMAN 18907 1.02 4.4E-01 AW080785.1 EST_HUMAN 18923 32818 0.89 4.4E-01 AW080785.1 EST_HUMAN 20325 <td>2432</td> <td></td> <td></td> <td>3.26</td> <td>4.4E-01</td> <td>P49765</td> <td>SWISSPROT</td> <td>VASCULAR ENDOTHELIAL GROWTH FACTOR B PRECURSOR (VEGF-B) (VEGF RELATED FACTOR)</td>	2432			3.26	4.4E-01	P49765	SWISSPROT	VASCULAR ENDOTHELIAL GROWTH FACTOR B PRECURSOR (VEGF-B) (VEGF RELATED FACTOR)
15965 28443 1.27 4.4E-01 AF058790.1 NT 15969 28446 2.31 4.4E-01 BF056728.1 EST_HUMAN 17706 2.07 4.4E-01 BE13396.1 EST_HUMAN 17839 30265 0.94 4.4E-01 BE141396.1 EST_HUMAN 17874 30692 4.06 4.4E-01 AW814885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW814885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW814885.1 EST_HUMAN 18505 31215 1.72 4.4E-01 AW814885.1 EST_HUMAN 18505 31215 1.52 4.4E-01 AV720408.1 INT 18724 31215 1.53 4.4E-01 AV720408.1 EST_HUMAN 18724 31477 1.53 4.4E-01 AV720408.1 EST_HUMAN 18905 31771 1.69 4.4E-01 AV720408.1 EST_HUMAN 18926 31477 1.53 4.4E-01 AV720408.1 EST_HUMAN 20325 32818 0.89 4.4E-01 AV720408.1 EST_HU	3357	1		1.27	4.4E-01		NT T	Rattus norvegicus SynGAP-b mRNA, complete cds
15969 28446 2.31 4.4E-01 BF058726.1 EST_HUMAN 1706 2.07 4.4E-01 BE378707.1 EST_HUMAN 1708 2.07 4.4E-01 BE378707.1 EST_HUMAN 17839 30265 0.94 4.4E-01 BE141396.1 EST_HUMAN 18242 30692 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 AW614885.1 EST_HUMAN 18505 31215 1.72 4.4E-01 SE5019.1 NT 18506 31215 1.72 4.4E-01 AV720408.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AV720408.1 EST_HUMAN 18724 31477 1.53 4.4E-01 AV080785.1 EST_HUMAN 18929 31771 1.69 4.4E-01 AV080785.1 EST_HUMAN 18923 32818 0.89 4.4E-01 AV080785.1 EST_HUMAN 20325 10.05 4.4E-01 AV080785.1 EST_HUMAN 21	3357	15965		1.27	4.4E-01	AF058790.1	NT	Rattus norvegicus SynGAP-b mRNA, complete cds
15804 1.28 4.4E-01 BE378707.1 EST_HUMAN 17706 2.07 4.4E-01 BE141398.1 EST_HUMAN 17839 30285 0.94 4.4E-01 U61154.1 NT 17874 0.9 4.4E-01 AW614885.1 EST_HUMAN 18242 30692 4.06 4.4E-01 AW614885.1 EST_HUMAN 18242 30693 4.06 4.4E-01 P04929 SWISSPROT 18489 31215 1.72 4.4E-01 S04029 SWISSPROT 18505 31231 1.9 4.4E-01 AV720408.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AV720408.1 EST_HUMAN 18724 31477 1.53 4.4E-01 AV198413.1 EST_HUMAN 18920 31771 1.69 4.4E-01 AV080785.1 EST_HUMAN 18923 32818 0.89 4.4E-01 AV080785.1 EST_HUMAN 20325 1.0.05 4.4E-01 AV080785.1 EST_HUMAN 21614 3450 0.76 4.4E-01 AV080780.1 NT 21647 3454	3361	15969			4.4E-01		EST HUMAN	7)91402.y1 NCI CGAP Br16 Homo sapiens cDNA clone IMAGE:3393795 5'
17706 2.07 4.4E-01 BE141396.1 EST_HUMAN 17839 30265 0.94 4.4E-01 U61154.1 NT 17874 0.9 4.4E-01 U61154.1 NT 18242 30692 4.06 4.4E-01 P04929 SWISSPROT 18242 30693 4.06 4.4E-01 P04929 SWISSPROT 18489 31215 1.72 4.4E-01 P04929 SWISSPROT 18505 31231 1.9 4.4E-01 AV720408.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AV198413.1 EST_HUMAN 18929 31771 1.69 4.4E-01 AV080785.1 EST_HUMAN 18920 31771 1.69 4.4E-01 AV080785.1 EST_HUMAN 20325 10.05 4.4E-01 AA080785.1 EST_HUMAN 21237 34160 1.01 4.4E-01 AA086427.1 EST_HUMAN 21647 34597 0.56 4.4E-01 AA086427.1 <t< td=""><td>4318</td><td>1</td><td></td><td>1.28</td><td></td><td></td><td>ĺ</td><td>601237139F1 NIH_MGC_44 Homo sapiens cDNA clone IMAGE:3609393 5</td></t<>	4318	1		1.28			ĺ	601237139F1 NIH_MGC_44 Homo sapiens cDNA clone IMAGE:3609393 5
17839 30265 0.94 4.4E-01 U61154.1 NT 17974 0.9 4.4E-01 AW614885.1 EST_HUMAN 18242 30682 4.06 4.4E-01 P04929 SWISSPROT 18242 30683 4.06 4.4E-01 P04929 SWISSPROT 18242 30683 4.06 4.4E-01 P04929 SWISSPROT 18505 31231 1.72 4.4E-01 S04029 SWISSPROT 18506 31215 1.72 4.4E-01 AV720408.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AV198413.1 EST_HUMAN 18920 31771 1.69 4.4E-01 AV080795.1 EST_HUMAN 20325 10.05 4.4E-01 AV080795.1 EST_HUMAN 21237 34160 1.01 4.4E-01 AV080795.1 EST_HUMAN 21647 34597 0.76 4.4E-01 AV080795.1 EST_HUMAN 21731 34674 1.01 4.4E-01 AV080795.1 EST_HUMAN 21647 34670 1.13 4.4E-01 AV080795.1 EST_HUMAN	5134	17706		2.07	4.4E-01	BE141396.1	!	MR0-HT0078-131289-007-005 HT0078 Homo sapiens cDNA
17874 0.9 4.4E-01 AW814885.1 EST_HUMAN 18242 30692 4.06 4.4E-01 P04929 SWISSPROT 18242 30693 4.06 4.4E-01 P04929 SWISSPROT 18489 31215 1.72 4.4E-01 S65019.1 NT 18724 31476 1.53 4.4E-01 AV720408.1 EST_HUMAN 18724 31477 1.53 4.4E-01 AI198413.1 EST_HUMAN 18929 31771 1.53 4.4E-01 AI198413.1 EST_HUMAN 19921 31771 1.69 4.4E-01 AV080795.1 EST_HUMAN 20325 10.05 4.4E-01 AA080795.1 EST_HUMAN 21237 34160 1.01 4.4E-01 AA080427.1 NT 21647 34549 0.76 4.4E-01 AF112540.1 NT 21647 34674 1.13 4.4E-01 AF112540.1 NT	5277	17839			4.4E-01	U61154.1		Buzura suppressaria nucleopolyhedrosisyirus ecdysteroid UDP-olucosytransferase (ent) gene complete cris
18242 30692 4.06 4.4E-01 P04929 SWISSPROT 18242 30693 4.06 4.4E-01 P04929 SWISSPROT 18489 31215 1.72 4.4E-01 S65019.1 NT 18724 31476 1.53 4.4E-01 A198413.1 EST_HUMAN 18724 31477 1.53 4.4E-01 A198413.1 EST_HUMAN 18929 31771 1.69 4.4E-01 ANO80795.1 EST_HUMAN 19921 32818 0.89 4.4E-01 ANO80795.1 INT 20325 10.05 4.4E-01 AA086427.1 INT 21637 3459 0.76 4.4E-01 AA086427.1 EST_HUMAN 21647 34597 0.76 4.4E-01 AA112540.1 NT 21731 34674 1.13 4.4E-01 AAF112540.1 NT	5417	17974		6:0	4.4E-01	AW814885.1	EST HUMAN	MR1-ST0206-120400-022-007 ST0208 Homo sapiens cDNA
18242 30693 4.06 4.4E-01 P04929 SWISSPROT 18489 31215 1.72 4.4E-01 S65019.1 NT 18724 31478 1.53 4.4E-01 A198413.1 EST_HUMAN 18724 31477 1.53 4.4E-01 A198413.1 EST_HUMAN 18980 31771 1.69 4.4E-01 ANO80795.1 EST_HUMAN 19971 1.02 4.4E-01 ANO80795.1 EST_HUMAN 20325 10.05 4.4E-01 ANO80795.1 NT 21237 34160 1.01 4.4E-01 ANO80795.1 NT 21614 34549 0.76 4.4E-01 ANO80427.1 NT 21637 34674 1.13 4.4E-01 ANG12578.1 EST_HUMAN 21731 34674 1.13 4.4E-01 ANG12578.1 EST_HUMAN	5613				4.4E-01		SWISSPROT	HISTIDINE-RICH GLYCOPROTEIN PRECURSOR
18489 31215 1,72 4,4E-01 S65019.1 NT 18505 31231 1,9 4,4E-01 AV720408.1 EST_HUMAN 18724 31476 1,53 4,4E-01 A198413.1 EST_HUMAN 18920 31771 1,69 4,4E-01 AN080795.1 EST_HUMAN 19071 1,02 4,4E-01 AN080795.1 EST_HUMAN 20325 32818 0,89 4,4E-01 AR080797.1 NT 21237 34160 1,01 4,4E-01 AN080797.1 NT 21647 34549 0,78 4,4E-01 AN08079.1 NT 21647 34674 1,13 4,4E-01 AN08079.1 NT	5613				4.4E-01		SWISSPROT	HISTIDINE-RICH GLYCOPROTEIN PRECURSOR
18724 31231 1.9 4.4E-01 AV720408.1 EST_HUMAN 18724 31476 1.53 4.4E-01 AI198413.1 EST_HUMAN 18930 31771 1.69 4.4E-01 AV080795.1 EST_HUMAN 19071 1.02 4.4E-01 AA776132.1 EST_HUMAN 20325 32818 0.89 4.4E-01 AA76132.1 EST_HUMAN 21237 34160 1.01 4.4E-01 AA056427.1 NT 21614 34549 0.78 4.4E-01 AA712540.1 NT 21647 34587 0.56 4.4E-01 AA712540.1 NT 21731 34674 1.13 4.4E-01 AAF112540.1 NT	5867				4.4E-01		N.	mucin frats, Sprague-Dawley, sulfur-dloxide-treated tracheal epithelium, mRNA Partial, 390 nt]
18724 31476 1.53 4.4E-01 AI198413.1 EST_HUMAN 18920 31771 1.69 4.4E-01 AW080795.1 EST_HUMAN 19071 1.02 4.4E-01 AW080795.1 EST_HUMAN 19953 32818 0.89 4.4E-01 AR000571.1 NT 20325 10.05 4.4E-01 AA056427.1 EST_HUMAN 21237 34160 1.01 4.4E-01 AA056427.1 EST_HUMAN 21614 34549 0.78 4.4E-01 AA112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 34587 1.13 4.4E-01 AW612578.1 EST_HUMAN	5883	18505		1.9	4.4E-01	AV720408.1	EST_HUMAN	AV720408 GLC Homo sapiens cDNA clone GLCCSC12 5'
18724 31477 1.53 4.4E-01 AI198413.1 EST_HUMAN 18980 31771 1.69 4.4E-01 AV080795.1 EST_HUMAN 19071 1.02 4.4E-01 AA76132.1 EST_HUMAN 19953 32818 0.89 4.4E-01 AA76132.1 EST_HUMAN 20325 10.05 4.4E-01 AA76132.1 INT 21237 34160 1.01 4.4E-01 AA056427.1 INT 21614 34549 0.76 4.4E-01 AA112540.1 INT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 - 34674 1.13 4.4E-01 OG2836 SWISSPROT	6108				4.4E-01	A1198413.1	EST_HUMAN	qi82h11.x1 NCI_CGAP_Bm25 Homo sepiens cDNA clone IMAGE:1881125 3' similer to TR:Q29168 Q29168 UNKNOWN PROTEIN
18980 31771 1.69 4.4E-01 AW080795.1 EST_HUMAN 19071 1.02 4.4E-01 AA776132.1 EST_HUMAN 19953 32818 0.89 4.4E-01 AA600571.1 NT 20325 10.05 4.4E-01 Z11679.1 NT 21237 34160 1.01 4.4E-01 AA056427.1 EST_HUMAN 21614 34549 0.76 4.4E-01 AA112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 - 34674 1.13 4.4E-01 O62836 SWISSPROT	6108	18724		1.53	4.4E-01	AI198413.1	EST_HUMAN	qi62h11.x1 NCI_CGAP_Brn25 Homo sepiens cDNA clone IMAGE:1861125 3' similar to TR:Q29168 Q29166 UNKNOWN PROTEIN ;
19071 1.02 4.4E-01 AZ78132.1 EST HUMAN 19953 32818 0.89 4.4E-01 AE00571.1 NT 20325 10.05 4.4E-01 Z11679.1 NT 21237 34160 1.01 4.4E-01 Z11679.1 NT 21614 34549 0.76 4.4E-01 AF112540.1 NT 21647 34587 0.56 4.4E-01 AM612578.1 EST HUMAN 21731 34674 1.13 4.4E-01 AM612578.1 EST HUMAN	6387	18990		1.69	4.4E-01	AW080795.1	EST_HUMAN	xc27e08.x1 NC_CGAP_Co18 Homo sepiens cDNA clone INAGE:2585510 3' similar to TR:085154 095154 AFLATOXIN B1-ALDEHYDE REDUCTASE.
19953 32818 0.89 4.4E-01 AE00571.1 NT 20325 10.05 4.4E-01 Z11679.1 NT 21237 34160 1.01 4.4E-01 AA056427.1 EST_HUMAN 21614 34549 0.76 4.4E-01 AF112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 -34674 1.13 4.4E-01 O62836 SWISSPROT	6470	19071		1.02	4.4E-01	AA776132.1	EST HUMAN	8885d11.s1 Stratagene schizo brain S11 Homo sapiens cDNA clone IMAGE:970965 3' similar to gb:M16038 TYROSINE-PROTEIN KINASE LYN (HUMAN):
20325 10.05 4.4E-01 Z11679.1 NT 21237 34160 1.01 4.4E-01 A056427.1 EST_HUMAN 21614 34549 0.76 4.4E-01 AF112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 - 34674 1.13 4.4E-01 O62836 SWISSPROT	7429	19953	32818		4.4E-01	AE000571.1	Į.	Helicobacter pylori 26995 section 49 of 134 of the complete genome
21237 34160 1.01 4.4E-01 AA056427.1 EST_HUMAN 21614 34549 0.76 4.4E-01 AF112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 334674 1.13 4.4E-01 O62836 SWISSPROT	7782	20325		10.05	4.4E-01		LN	S.tuberosum mRNA for induced stoken tip protein (partial)
21614 34549 0.76 4.4E-01 AF112540.1 NT 21647 34587 0.56 4.4E-01 AW612578.1 EST_HUMAN 21731 . 34674 1.13 4.4E-01 O62836 SWISSPROT	8698	21237	34160	1.01	4.4E-01	AA056427.1	EST_HUMAN	z69a03.s1 Stratagene colon (#637204) Homo sapiens cDNA clone IMAGE:508836 3'
21647 34587 0.56 4.4E.01 AW612578.1 EST_HUMAN 21731 3.4674 1.13 4.4E.01 062836 SWISSPROT	8078	21614	34549	92.0	4.4E-01	AF112540.1	NT	HIV-1 isolate 08107v6 from USA, envelope glycoprotein (env) gene, partial cds
21731 34674 1.13 4.4E-01 062836 SWISSPROT	9111	21647		0.56			EST_HUMAN	hh05c08.x1 NCI_CGAP_Kid11 Hamo sapiens cDNA clone IMAGE:2854222 3' similar to SW:MSH6_HUMAN P52701 DNA MISMATCH REPAIR PROTEIN MSH6 :
	9214	21731	Ы	1.13	4.4E-01		SWISSPROT	ZINC FINGER X-CHROMOSOMAL PROTEIN

Page 56 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

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Top Hit Descriptor	qo39f09.x1 NCI CGAP Lu5 Homo sabiens cDNA clone IMAGE-1910821 3	GLYCOPROTEIN B PRECURSOR (GLYCOPROTEIN 14)	TYROSINE-PROTEIN KINASE RECEPTOR TIE-1 PRECURSOR	beta -HKA=H,K-ATPase beta-subunit frats. Genomic. 8983 nt. segment 2 of 21	beta -HKA=H,K-ATPase beta-subunit frats, Genomic, 8963 nt, segment 2 of 2]	Mus musculus sodium channel, type X, alpha polypeotide (Scn10a), mRNA	Homo sepiens chromosome 21 segment HS21C082	UV EXCISION REPAIR PROTEIN PROTEIN RADZ3 HOMOLOG A (HHR23A)	Callithrix Jacchus MW/LW opsin gene, upstream flanking region	Callithrix jacchus MW/LW opsin gene, upstream flanking region	MR0-BN0070-270300-008-g04 BN0070 Homo sapiens cDNA	Human somatostatin I gene and flanks	Callithrix jacchus MW/LW opsin gene, upstream flanking region	Callithrix jacchus MW/LW opsin gene, upstream flanking region	LARGE PROLINE-RICH PROTEIN BAT2 (HLA-B-ASSOCIATED TRANSCRIPT 2)	LARGE PROLINE-RICH PROTEIN BAT2 (HLA-B-ASSOCIATED TRANSCRIPT 2)	QV1-HT0638-070500-191-d08 HT0638 Homo sapiens cDNA	Saimiri sclureus offactory receptor (SSC188) gene, partial ods	Coturnix coturnix Japonica (inG gene	DNA GYRASE SUBUNIT B	602023134F1 NCI_CGAP_Brn67 Homo sapiens cDNA clone IMAGE:4158296 5'	Methanococcus voltae flagella-related protein C-I (flaC-flal) genes, complete cds	Erwinia amylovora rcsV gene	hh74e10.y1 NCI_CGAP_GU1 Homo sapiens cDNA clone IMAGE.2968554 5'	hh74e10.y1 NCI_CGAP_GU1 Hamo sepiens cDNA clone IMAGE.2968554 5'	хи83e05.x1 Soares_NHCeC_cervical_tumor Homo sapiens cDNA clone IMAGE.2698400 3' similar to TR-000189 000189 Mil. ADAPTIN. PET ATED PROTEIN 2	Equus caballus microsabellite LEXO27	284404 x 1 NCI CGAP Ov35 Homo saniens cDNA clone IMAGE 2293351 31	LARGE PROLINE-RICH PROTEIN BAT2 (HLA-B-ASSOCIATED TRANSCRIPT 2)	LARGE PROLINE-RICH PROTEIN BAT2 (HLA-B-ASSOCIATED TRANSCRIPT 2)	Streptomyces coelicolor whill gene	CELL DIVISION PROTEIN FTSH HOMOLOG PRECURSOR	nz24e09.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1288696 3'
Top Hit Database Source	EST HUMAN	SWISSPROT	SWISSPROT	LN	LN	N _T	LN	SWISSPROT	N	FZ	EST_HUMAN	۲	LN.	LN	SWISSPROT	SWISSPROT	EST HUMAN	LΝ	ΤN	SWISSPROT	EST_HUMAN	NT	IN	EST_HUMAN	EST_HUMAN		Ξ	EST HUMAN	SWISSPROT	SWISSPROT	Į.	SWISSPROT	EST_HUMAN
Top Hit Acession No.	AI268650.1	P28922	P35590	S76404.1	S76404.1	6877874 NT	AL163282.2	P54725	AF155218.1	AF155218.1	AW999477.1	J00306.1	AF155218.1	AF155218.1	P48634	P48634	BE181655.1	AF179825.1	AJ001678.1	033367	BF348001.1	U97040.1	Y14604.1	AW630048.1	AW630048.1	AW170559.1	AF075629.1	AI874332.1	P48634	P48634	AJ003022.1	Q39102	AA761653.1
Most Similar (Top) Hit BLAST E Value	4.4E-01	_	4.4E-01	4.4E-01	4.4E-01	4.4E-01	4.4E-01	4.4E-01			4.3E-01	4.3E-01		4.3E-01	4.3E-01				4.3E-01	4.3E-01	4.3E-01			ļ	4.3E-01	4.3E-01	_			4.3E-01	4.3E-01		4.2E-01
Expression Signal	1.69	2.12	4.51	1.43	1.43	4.68	14.98	1.5	1.77	1.77	0.91	1.21	3.96	3.96	0.76	0.78	1.34	2.08	4.28	0.78	1.76	2.66	0.7	2.63	2.63	0.57	2.52	13.	1.55	1.55	2.81	1.39	1.04
ORF SEQ ID NO:	35347		35495	65758	35760	31016				25565				25565					32215				34702	35109	35110	35614	32451	37101	30645	30646		26524	
SEQ ID	22369	22370	22504		22771	24271	25000	24635	- 1		15711	16819				18198	1	- 1	J	İ		20806	21758	22142	22142	22623	19618	24031	18198	18198	24696	15440	14573
Probe SEQ ID NO:	9872	9873	10009	10276	10276	11939	11952	12517	438	436	3096	4231	4495	4495	5587	5567	6049	6065	88	6949	7456	8366	9179	8642	9642	10128	10811	11588	11632	11632	12616	1402	1991

Page 57 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

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Probe SEQ ID NO:	_ v	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
2068			1.37	4.2E-01	AF258325.1	Þ	Plasmodium falciparum multidrug resistance protein Pgh1 gene, complete cds
3669			4.91	4.2E-01	AE003947.1	F	Xyiella fastidiosa, section 93 of 229 of the complete genome
3699		28768	1	4.2E-01	A1280338.1	EST_HUMAN	q94b01 x1 Soares_NhHMPu_S1 Homo sapiens cDNA clone IMAGE:1876945 3
3773	18014		9.0	4.2E-01	N81203.1	EST_HUMAN	788iE1 fetal brain cONA Homo sapiens cONA clone 788iE1-K similar to R07879, Z40498
3948	16546	29014	0.73	4.2E-01	AW 835527.1	EST_HUMAN	QV0-LT0015-180200-127-h01 LT0015 Homo sapiens cDNA
4054	16651		86.0	4.2E-01	Q04886	SWISSPROT	SOX-8 PROTEIN
4807	17385	29835	4.3	4.2E-01	AA534083.1	EST HUMAN	nj69h01.s1 NCI_CGAP_P10 Homo sapiens cDNA done IMAGE:997777 similar to gb:M33600 HLA CLASS II HISTOCOMPATIBILITY ANTIGEN DR-1 RETA CHAIN (HIMAN)
4895	17470		4	4.2E-01		EST HUMAN	V77601.r1 Soares Infant brain 1NIB Homo sapiens CONA clone IMAGE-28278.5'
5232	17796		3.77	4.2E-01	U50871.1	IN	Human familial Alzheimer's disease (STM2) gene, complete cds
5891	18514	31241	1.52	4.2E-01	BF242055.1	EST HUMAN	601879721F1 NIH MGC 55 Homo sepiens cDNA clone IMAGE:4108493 5
5953	18575	31309	2.16	4.2E-01	AW854162.1	EST_HUMAN	RC3-CT0254-060400-028-g04 CT0254 Homo sapiens cDNA
6352			1.08	4.2E-01	AL163247.2	-N	Homo sapiens chromosome 21 segment HS21C047
7031	19565		62.01	4.2E-01	AU158472.1	EST_HUMAN	AU158472 PLACE2 Homo sapiens cDNA clone PLACE2000470 3'
7031	19565	32383	10.29	4.2E-01	AU158472.1	EST_HUMAN	AU158472 PLACE2 Homo sapiens cDNA clone PLACE2000470 3'
7082			1.97	4.2€-01	\$82504.1	ĖΝ	Brca1=breast cancer gene [rats, WF, splean, Genomic, 419 nt, segment 2 of 2]
7150			5.81	4.2E-01	AL161547.2	ΤN	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 47
7934						EST_HUMAN	EST369413 MAGE resequences, MAGE Homo sapiens cDNA
7934	20478	33388	2.61	4.2E-01	AW957448.1	EST_HUMAN	EST369413 MAGE resequences, MAGE Homo sapiens cDNA
8148	20889	33602	0.55	4.2E-01	4758039 NT	Į.	Homo sapiens cytochrome c oxidase subunit Vic (COX6C), nuclear gene encoding mitochondrial protein, mRNA
9235				4.2E-01	U57431.1	FN	Human cytomegalovirus early phosphoprotein p50 mRNA, complete cds
9235		34707	0.52	4.2E-01	U57431.1	NT	Human cytomegalovirus early phosphoprotein p50 mRNA, complete cds
9880			0.81	4.2E-01		EST_HUMAN	295f01.s1 Soares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA clone IMAGE:462649 3'
10083						NT	Lassa virus strain 803213 glycoprotein precursor and nucleoprotein genes, complete cds
10390						EST_HUMAN	MR3-SN0010-280300-103-h07 SN0010 Homo sapiens cDNA
10921			3.69	4.2E-01		NT	Oryzias latipes OIGC7 mRNA for membrane guanyly cyclase, complete cds
11273		36780	2.65		BE966485.2	EST_HUMAN	801680352R1 NIH_MGC_71 Hamo sapiens cDNA clane IMAGE:3906085 3'
12561			1.49	4.2E-01	AV731815.1	EST_HUMAN	AV731815 HTF Homo sapiens cDNA clone HTFBHH05 5'
1133			1.59	4.1E-01	Al905481.1	EST_HUMAN	RC-BT091-210199-142 BT091 Homo sapiens cDNA
1142	1		1.54	4.1E-01	AV705243.1	EST_HUMAN	AV705243 ADB Homo sapiens cDNA clone ADBAHF08 5'
1142			2.52	_	AV705243.1	EST_HUMAN	AV705243 ADB Homo sapiens cDNA clone ADBAHF08 5'
2735	J			4.1E-01	5283	L	Homo sapiens anaphase-promoting complex subunit 7 (APC7), mRNA
2967	15582	78061	2.11	4.1E-01	AL161536.2	N	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 36

Page 58 of 526 Table 4 Single Exon Probes Expressed in Fetal ∟lver

Top Hit Descriptor	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 36	ol94b08.s1 Scares_NFL_T GBC_S1 Homo saplens cDNA clone IMAGE:1505943.3	EST373364 MAGE resequences, MAGG Homo sapiens cDNA	EST373384 MAGE resequences, MAGG Homo sapiens cDNA	Rhodococcus sp. AD45 isoG, isoH, Isol, isoJ, isoA, isoB, IsoC, IsoD, IsoE and isoF genes	om33d02.s1 Sceres_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:1542819.37	AV747880 NPC Homo sepiens cDNA clone NPCBDF10 5'	602156590F1 NIH_MGC_83 Homo sapiens cDNA clone IMAGE:4297319 5	Methanococcus jannaschii section 77 of 150 of the complete genome	602133261F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4288238 5'	Mus musculus signaling intermediate in Toll pathway-evolutionarily conserved (Sitpec-pending). mRNA	Voalavo gymnocaudus Vgym560 cytochrome b (cytb) gene, complete cds; mitochondrial gene for	mitochondrial product	Cempylobacter jejuni NCTC11168 complete genome; segment 3/6	AV649579 GLC Homo sapiens cDNA clone GLCBVD123'	PROBABLE SERINE PROTEASE DO-LIKE PRECURSOR (59 KDA IMMUNOGENIC PROTEIN) (SK59)	PROBABLE SERINE PROTEASE DO-LIKE PRECURSOR (59 KDA IMMUNOGENIC PROTEIN) (SK 59)	CM2-HT0137-200999-010-e08 HT0137 Homo sapiens cDNA	Zea mays ZMPMS2 gene for 19 kDa zein protein	VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KV1.1 (HUKI) (HBK1)	Homo sapiens DNA for amyloid precursor protein, complete cds	RC2-CT0201-290999-012-d10 CT0201 Homo sapiens cDNA	Laqueus rubellus mitochondrion, complete genome	Drosophila metanogaster Dalmatian (dmt) mRNA, complete cds	Mus musculus platelet derived growth factor receptor, beta polypeptide (Pdgfrb), mRNA	Ascobolus immersus masc2 gene	Ascobolus immersus masc2 gene	Deinococcus radiodurans R1 section 68 of 229 of the complete chromosome 1	Deinococcus radiodurans R1 section 68 of 229 of the complete chromosome 1	Mus musculus ubiquitin-protein ligase 83 componen n-recognin (Ubr1), mRNA	Homo sapiens chromosome 21 segment HS21 C080	Homo sepiens chromosome 21 segment HS21 0080
Top Hit Database Source	¥	T HUMAN	EST HUMAN	EST HUMAN	Т	EST HUMAN	EST HUMAN		N L	EST_HUMAN			NT	N	EST_HUMAN	SWISSPROT	SWISSPROT	EST_HUMAN	Ī	SWISSPROT	INT	EST_HUMAN		I LN		NT	LN	Į.	- IN			IN
Top Hit Acession No.	AL161536.2	AA906344.1	AW961292.1	AW961292.1	AJ249207.1	AA909257.1	AV747880.1	BF681393.1	U67535.1	BF574604.1	6755521 NT			AL139076.2	AV649579.1	P18584	P18584	BF349382.1	X58700.1	Q09470		AW847123.1	8404656 NT	AF203478.1	6679258 NT	Z96933.1	296933.1	AE001931.1	AE001831.1	6678490 NT		AL 163280.2
Most Similar (Top) Hit BLAST E Value	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01		4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.1E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01	4.0E-01
Expression Signal	2.11	99.0	0.58	0.58	2.82	0.76	1.31	3.97	2.74	1.31	1.26		0.61	1.26	0.79	0.51	0.51	2.29	45.22	3.57	2.6	4.55	0.82	1.51	4.1	1.22	1.22	17.82	17.82	1.45	1.23	1.23
ORF SEQ ID NO:	28062	28428	28899	28900	29390		28807	31513	32848	33427	34484						35891		36262	36015				26505				27352	27353	25299	28090	28091
Exen SEQ ID NO:	15582	15952	16438	16438	16948	16979	17355	18755		20521	21556	- !	21941			22895	22895		23247			15408	13681	13978	- 1		- 1	14780				15611
Probe SEQ ID NO:	2967	3342	3839	3839	4361	4393	4774	6141	7460	7979	9019		<u>8</u>	10164	10310	10401	10401	10471	10719	11270	12290	147	1077	138	1532	2049	2049	2204	2204	2831	2885	2895

Page 59 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	Exan SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
3758	16359	28829	2.17	4.0E-01	AF068903.1	<u> </u>	Streptococcus pneumoniae YIIC (yIIC), YIID (yIID), penicillin-binding protein 2x (pbp2x), and undecaprenyiphosphate-UDP-MurNAc-pentapeptide phospho-MurNAc-pentapeptide transferase (mraY) genes, complete cds.
3899	16498				AJ277511.1	Ę	Ovis aries partial JD2 gene for T cell receptor delta chain (TCRDJ2), exon 1
3899	16498		3.04	4.0E-01	AJ277511.1	Z	Ovis aries partial JD2 gene for T cell receptor delta chain (TCRDJ2), exon 1
4942	17517		8.41	4.0E-01	Q31849	SWISSPROT	NADH-PLASTOQUINONE OXIDOREDUCTASE CHAIN 5, CHLOROPLAST
6909	18686	31429	1.16	4.0E-01	AW970610.1	EST_HUMAN	EST382691 MAGE resequences, MAGK Homo sapiens cDNA
6567	19165	31961	78.0	4.0E-01	P27285	SWISSPROT	STRUCTURAL POLYPROTEIN (P130) [CONTAINS: COAT PROTEIN C; SPIKE GLYCOPROTEINS E3, E2 AND E1; 8 KD PEPTIDE]
7777	20286	33183	0.72	4.0E-01	P27548	SWISSPROT	MICROTUBULE ASSOCIATED PROTEIN 4
7869	20411	33317	0.48	4.0E-01	BF092634.1	EST_HUMAN	MR4-TN0110-180900-202-g02 TN0110 Home saplens cDNA
7954	20496	33408	0.99	4.0E-01	AB016625.1	Z	Homo sapiens OCTN2 gene, complete cds
9839	21474	34394	96.0	4.0E-01	AA323289.1	EST_HUMAN	EST26066 Cerebellum II Homo sepiens cDNA 5' end similar to EST containing Alu repeat
11443	23883		1.65	4.0E-01	BF030262.1	EST_HUMAN	601558283F1 NIH_MGC_58 Homo sepiens cDNA clane IMAGE:3828092 5'
11568	24015		3.52	4.0E-01	L76080.1	NT	Synechocystis sp. PCC 9413 transposase gene, complete cds
11958	24901		2.5	4.0E-01	AL163300.2	F	Homo sapiens chromosome 21 segment HS21C100
12518	24636		1.42	4.0E-01	P36049	SWISSPROT	HYPOTHETICAL 49.7 KD PROTEIN IN GINZ-STE3 INTERGENIC REGION
1420	14013	26543	1.98	3.9E-01	AF206618.1	MT	Gorilla gorilla carboxyl-ester lipase (CEL) gene, complete cds
2668					AB033019.1	NT	Homo sapiens mRNA for KIAA1193 protein, partial cds
2730			3.79			NT	H.saplens B-myb gene
2730					X82032.1	NT	H.sapiens B-myb gene
3131			3.95		AJ225896.1	NT	Sinorhizoblum mellioti egi, syrB2, cya3 genes and orf3
4153	16745	29199	1.49	3.9E-01	BF592611.1	EST_HUMAN	7/61401.x1 NCI_CGAP_Br16 Homo sapiens cDNA clone IMAGE:3339169 3'
5130	17702	30136	1.86	3.9E-01	BE728667.1	EST_HUMAN	601563948F1 NIH_MGC_20 Homo seplens cDNA clone IMAGE:3833699 5'
0609	18708	31454	6.44	3.9E-01	BF208036.1	EST_HUMAN	601862362F1 NIH_MGC_53 Homo saplens cDNA done IMAGE:4082055 5'
8438	19030	31812	890	3 05-04	c soscali	<u>F</u>	Homo saplens zinc finger protein 92 (ZFP92), expressed-Xq283TS protein (XQ280RF), and biglycan (BGN)
7896				L	U79415.1	Į.	Homo seplens preprior dipeopledy peopled to PPP-I) sense complete cds
8795	1					EST HUMAN	CM3-CT0105-170899-004-b08 CT0105 Homo sapiens cDNA
8804	21343		0.7	3.9E-01	BF348634.1	EST_HUMAN	602019844F1 NCI_CGAP_Brn67 Homo sapiens cDNA clone IMAGE:4155322 5
9161		34640	1.24	3.9E-01	AW 195888.1	EST HUMAN	xn88d04.x1 Sogres_NFL_T_GBC_S1 Home sapiens cDNA clone IMAGE:2701351 3' similar to TR:094821 C094821 KIAA0713 PROTEIN ;
	J	l					

Page 60 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	_	_			_						_						_					_				_	_		_			
Top Hit Descriptor	wp78a02x1 NCI_CGAP_Bm25 Homo sapiens cDNA clone INAGE:2467658 3' similar to SW:RFX5 HUMAN P48382 BINDING REGULATORY FACTOR	Human clabindin 27 gene, exons 10 and 11, and L1 and Alu repeats	Porphyra purpurea mitochondrion, complete genome	Nicotiana tabacum mRNA for TATA binding protein (TBP), complete cds	Human beta-B2-cnystallin (B2-1) gene, exon 4, partial cds	AV695974 GKC Hamo sapiens cDNA clone GKCBQC11 5	Homo sapiens proteoglycan 3 (PRG3) gene, complete cds	HOMEOBOX PROTEIN HLX1	Thermotoga maritima section 123 of 136 of the complete genome	Homo sapiens hypothetical protein FLJ10583 (FLJ10583), mRNA	Homo sapiens protein kinase PKNbeta (pknbeta), mRNA	Mus musculus pcm-1 mRNA for percentriolar material-1, complete cds	Xyletla fastidiosa, section 16 of 229 of the complete genome	Arabidopsis thaliana putative c-myb-like transcription factor (MYB3R-3) mRNA complete cds	Mus musculus solute carrier family 1, member 6 (Slc1a6), mRNA	Human immunodeficiency virus type 1 complete genome (isolate 98SE-MP1213)	Pieuronectes americanus aminopeptidase N (ampN) gene, partial cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 30	w/38b12.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2357855 3'	w/38b12.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2357855 3:	PMO-HT0339-200400-010-G01 HT0339 Homo sapiens cDNA	Mus musculus general transcription factor II I (Gtf2), mRNA	Takifugu rubripes wnt2 (partial), frank1, cftr and frank2 (partial) genes	601074110F1 NIH_MGC_12 Homo sapiens cDNA clone IMAGE:3460154 5'	yr68a11.r1 Sogres fetal liver spleen 1NFLS Hamo sapiens cDNA clone IMAGE:210428 5' similar to	gb M87933 HUMAALU364 Human carcinoma cell-derived Alu RNA transcript, (rRNA); gb:M96956	EPIDERMAL GROW TH FACTOR-LIKE CRIPTO PROTEIN (HUMAN); contains Alu repetitive	element, contains MER4 repetitive element :	TRANSCRIPTION FACTOR SOX-10	prion protein [mink, Genomic, 2448 nt]	QV3-BT0537-271299-049-e02 BT0537 Homo sapiens cDNA	ta54f11.x1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone IMAGE:2047917.3 similar to contains Atu repetitive element.
Top Hit Defeabase Source	EST HUMAN	N	ĽΝ	<u>⊼</u>	١	EST HUMAN	N	SWISSPROT	N.	Ä	۲	Ę	Į.	Į.	LN L	۲	Ę	ᅜ	EST_HUMAN	EST_HUMAN	EST_HUMAN	IN	NT	EST_HUMAN				EST_HUMAN	SWISSPROT	IN	EST_HUMAN	ESŤ_HUMAN
Top Hit Acession No.	A1937337.1	M19879.1	11465620 NT	D86722.1	M18440.1	AV695974.1	AF304354.1	Q61670	AE001811.1	11433335 NT	7019488 NT	AB029291.1	AE003870.1	AF214117.1	6678002 NT	AJ251057.1	AF043383.1	AL161518.2	Al807219.1	AI807219.1	BE154080.1	6754095 NT	AJ271361.2	BE544653.1				H64927.1	Q04888	S46825.1	BE072399.1	Al374601.1
Most Similar (Top) Hit BLAST E Vælue	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.9E-01	3.8E-01	3.8E-01	3.8E-01		3.8E-01	3.8E-01				3.8E-01	3.8E-01			3.8E-01	-						3.8E-01	3.8E-01
Expression Signal	1.42	3.68	0.5	0.69	0.48	1.82	3.42	1.42	1.56	1.37	19.28	3.11	0.99	1.89	3.94	0.89	2.2	9.83	0.59	0.75	0.94	0.8	0.69	86°C			•	1.07	1.11	0.68	5.29	3.97
ORF SEQ ID NO:	34945	35274		35561	25833				31015					27734	27791			28615			28882	29058	29183	30259		-		30378	31135		32137	32423
Exon SEQ ID NO:				22566	22986	23229		24214			12834	13162					15699			ı			16730	17833	-			- 1	18419	19082	19331	19591
Probe SEQ ID NO:	9464	8792	9828	10071	10492	10700	11729	11854	11930	12389	171	531	1911	2605	2881	3034	3084	3530	3592	3609	3820	3989	4138	5271				5412	5794	98	6737	6857

Page 61 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

			sp				ne IMAGE:120539 5' similar to contains			GE:30289 3'	GE:30289 3'				alete cds			mplete cds		JA clone IMAGE:15101883		the complete genome	spi		e cds			ANS.	ANA	16701 3'	(9)		MAGE:1950997 3'	1 and KIAA0851 gene
Top Hit Descriptor	Arabidopsis thaliana DNA chromosome 4, conlig fragment No. 25	M.musculus gene for kalilkrein-binding protein	Mouse liver receptor homologous protein (LRH-1) mRNA, complete cds	Homo sepiens mRNA for KIAA1631 protein, partial cds	Homo sapiens FOS-like antigen-1 (FOSL1), mRNA	Homo sapiens chromosome 21 segment HS21C079	ye43h06.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:120539 5' similar to contains	Alu repetitive element;contains PTK5 repetitive element;	RC0-HT0841-040800-032-b12 HT0841 Homo sapiens cDNA	y/92h11.s1 Soares infant brain 1NIB Homo sapiens cDNA clone IMAGE:30289 3'	y/92h11.s1 Sogres infant brain 1NIB Homo sapiens cDNA clone IMAGE:30289 3'	Borrelia burgdorferi (section 10 of 70) of the complete genome	Human p53 (TP53) gene, complete cds	QV3-ET0063-190700-271-e05 ET0063 Homo sepiens cDNA	Mus musculus apoptosis inhibitor bcl-x (bcl-x) gene, expn 3 and complete cds	Mus musculus developmental control protein mRNA, partial cds	Homo sapiens mRNA for KIAA1410 protein, partial cds	Danio rerio bone morphogenetic protein 4 precursor (BMP4) gene, complete cds	EST21715 Adrenal gland tumor Homo sapiens cDNA 5' end	ok39c07.x1 Sogres_NSF_F8_9W_OT_PA_P_S1 Homo sapiens cDNA clone IMAGE:1510188 3	MR3-OT0007-080300-104-b02 OT0007 Homo sapiens cDNA	Neisseria meningitidis serogroup B strain MC58 section 50 of 206 of the complete genome	Homo sapiens interferon-induced protein p78 (MX1) gene, complete cds	Homo sapiens chromosome 21 segment HS21C078	Chicken (White leghorn) delta-1 and delta-2 crystallin genes, complete cds	Mus sextode haptoglobin mRNA, complete cds	Homo saplens tumor endothelial marker 7 precursor (TEM7), mRNA	Homo sapiens chromosome 12 open reading frame 4 (C120RF4), mRNA	Homo sapiens chromosome 12 open reading frame 4 (C120RF4), mRNA	ok43b11.s1 NCI_CGAP_Lei2 Homo sapiens cDNA clone IMAGE:1518701 3'	Gallus gallus mRNA for beta-carotene 15,15'-dioxygenase (bCDO gene)	mouse ig germline alpha membrane exons region	qt46b07.x1 Soares_fetal_lung_NbHL19W Homo sapiens cDNA clone IMAGE:1950997 3'	Homo sapiens partial LIMD1 gene for LIM domains containing protein 1 and KIAA0851 gene
Top Hit Database Source	TN	۲	ΙN	Ę	N	NT		T			EST_HUMAN	LΝ	Į,	EST_HUMAN	Ł	Z	NT	LZ LZ	EST_HUMAN	EST_HUMAN	EST_HUMAN	ΝŢ	. LN	LN	TN	N			١	T_HUMAN	TN	LΝ	L_HUMAN	
Top Hit Acession No.	AL 161513.2	X61597.1	M81385.1	AB046851.1	11441284 NT	AL163279.2		195413.1	BE719219.1	R42550.1	R42550.1	AE001124.1	U94788.1	BE829256.1	U78031.1	AF194972.1	AB037831.1	AF056336.1	AA319482.1	A1218707.1	AW878037.1	AE002408.1	AF135187.1	AL163278.2	M10806.1	L10353.1	11525843 NT	11436739 NT	11436739 NT	AA902912.1	AJ271386.1	K00691.1	Al336411.1	AJ297357.1
Most Similar (Top) Hit BLAST E Value	3.8E-01/	_		3.8E-01	3.8E-01	3.8E-01						3.8E-01	3.8E-01	3.8E-01			3.7E-01			3.7E-01	3.7E-01	3.7E-01	3.7E-01				3.7E-01	3.7E-01	3.7E-01		3.7E-01	3.7E-01	3.7E-01	3.7E-01
Expression Signal	1.33	4.75	0.49	2.34	1.14	1.12		\$.U3	3.5	2.95	2.95	2.81	1.75	1.45	2.22	1.25	15.01	10.94	0.68	9.19	1.18	3,13	1.27	0.94	1	0.81	4.4	1.88	1.88	0.78	1.54	0.46	4.17	3.47
ORF SEQ ID NO:	32339			33951	34019	34210				37060					\mathbf{I}	30866	27657				29440	29514	31285	31474	32033		32579			33770			`.	38448
Exan SEQ ID NO:	19517			21031	21099	21290		┙		┙	┙	24272		24421			15085				16997	17064				19247	19728		L					23428
Probe SEQ ID NO:	7019	7525	8238	8492	8560	8751	90,0	3	11408	11541	11541	11940	12069	12189	12565	12638	2521	3507	3838	4313	4412	4479	5936	6105	9633	6651	7197	8271	8271	8308	9129	10074	10111	10909

Page 62 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	,	-	_	_	_	~	_	_	_	_	_	_	_	-	-	_		_	_		_		_		_		-	_	_	_	_
Top Hit Descriptor	Homo sapiens partial LIMD1 gene for LIM domains containing protein 1 and KIAA0851 gene	Bowine mRNA for terminal deoxynucleotidy/transferase (TdT) (EC 2.7.7.31)	0048d03.s1 NCI_CGAP_Lu5 Homo sapiens cDNA clone IMAGE:1569221.3' similar to gb:M77698 TRANSCRIPTIONAL REPRESSOR PROTEIN YY1 (HUMAN);	Mus musculus retinoblastoma 1 (Rb1), mRNA	Human heart/skeletal muscle ATP/ADP translocator (ANT1) gene, complete cds	Chlamydophila psittaci partial omp1 gene for cuter membrane protein 1	Human mRNA for KIAA0223 gene, partial cds	DKFZp762K075_r1 762 (synonym: hmel2) Homo sapiens cDNA clone DKFZp762K075 5'	Homo sapiens NF2 gene	Brassica napus mRNA for MAP4K atpha2 protein	Human mibp gene, partial cds	yd03e05.r1 Soares infant brain 1NIB Homo sapiens cDNA clone IMAGE: 24443 5'	yd03e05.r1 Soares infant brain 1NIB Homo saplens cDNA clone IMAGE;24443 5'	hg33f02.xf NCI_CGAP_GC6 Homo sepiens cDNA clone IMAGE.2947419 3'	hg33f02.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2947419 3'	Mus musculus ribosomal protein S19 (Rps19) gene, complete cds	Raftus norvegicus repeat element associated with the Rasgrf1 gene	Human mRNA for KIAA0323 gene, partial cds	P. Irregulare (P3804) gene for actin	RC5-ST0171-181099-011-g07 ST0171 Hamo sapiens cDNA	PROTEIN-L-ISOASPARTATE O-METHYLTRANSFERASE (PROTEIN-BETA-ASPARTATE	METHYLTRANSFERASE) (PIMT) (PROTEIN L-ISOASPARTYL METHYLTRANSFERASE) (L- ISOASDADTYL DROTEIN CADBOXYL METHYLTDANISEEDASE)	Operating melanopaster supertransporter 3 (sur3) mRNA complete of	H. sapiens serotonin transporter gene, exons 9 and 10	Hisablens serotonin transporter gene, exons 9 and 10	RC1-HT0545-150600-014-b12 HT0545 Homo sapiens cDNA	Brassica napus mRNA for MAP4K alpha2 protein	Z.mays mRNA for casein kinase II alpha subunit	ha02g04.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE:2872568 3'	MR4-BT0358-270300-005-c10 BT0358 Homo sapiens cDNA	FORMATE HYDROGENLYASE SUBUNIT 5 PRECURSOR (FHL SUBUNIT 5) (HYDROGENASE-3 COMPONENT E)
Top Hit Database Source	NT	IN	EST_HUMAN	NT	LN	ΝT	Į.	EST_HUMAN	TN	LΝ	Z	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	NT	TN	Ā	LN	EST HUMAN		TOddaalwa	TN	L	LN	EST HUMAN	LN	LN	EST_HUMAN	EST_HUMAN	SWISSPROT
Top Hit Acession No.	3.7E-01 AJ297357.1	X04122.1	3.7E-01 AA973540.1	TN 8477678		3.7E-01 AJ243525.1	3.7E-01 D86976.1	3.7E-01 AL121154.1	Y18000.1	3.6E-01 AJ009609.1	3.6E-01 U89241.1	T80255.1	T80255.1	3.6E-01 AW590184.1	4W590184.1	3.6E-01 AF216207.1	3.6E-01 AF056927.1	3.6E-01 AB002321.1	X76725.1	3.6E-01 AW812033.1		004006	3.6F-01 AF199485 1	X76758.1	X76758.1	3.6E-01 BE707883.1	3.6E-01 AJ009609.1	Y11526.1	3.6E-01 AW339393.1	3.6E-01 BE067699.1	P16431
Most Similar (Top) Hit BLAST E Value	3.7E-01	3.7E-01 X04122.1	3.7E-01	3.7E-01	3.7E-01 J04982.1	3.7E-01	3.7E-01	3.7E-01	3.7E-01 Y18000.1	3.6E-01	3.6E-01	3.6E-01 T80255.1	3.6E-01 T80255.1	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01 X76725.1	3.6E-01		3 GE_01 D2420G	3.6F-01	3.6E-01 X76758.1	3.6E-01 X76758.1	3.65-01	3.6E-01	3.6E-01 Y11526.1	3.6E-01	3.6€-01	3.6E-01 P16431
Expression Signal	3.47	4.81	1.6	2.78	1.82	4.15	4.72	2.94	7.01	1.07	8.45	4.32	4.32	6.39	6.39	7.23	0.88	1.13	2.49	3.34		4,	10.38	2.16	2.16	1.97	46.0	0.65	2.28	0.58	1.16
ORF SEQ ID NO:		36048				-			30904	52454		26477	26478	27099	27100					27665		27780		28600	28601	29530			30154	30249	31616
Exen SEQ ID NO:	23428	23039	23997	24046	24601	24191	24251	24499	24545	12938	13643	13951	13951	14543	14543	14576	14677	14881	15000	15092		15218	18012	16121	16121	17081	17428	17443	17723	17824	18845
Probe SEQ ID NO:	10909	11341	11549	11603	11640	11821	11913	12302	12377	282	1033	1357	1357	1959	1959	1994	2098	2309	2433	2528		2850	2924	3516	3516	4497	4850	4887	5153	5261	6236

WO 01/57277

Page 63 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

		-	_						_				_					_		_	-					_	_		
	Top Hit Descriptor	Homo sapiens PHEX gene	y74a06.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:275987 5'	wt72c10.x1 Soares_thymus_NHFTh Homo sapiens cDNA clone IMAGE:2513010 3' similar to TR:015117 015117 FYN BINDING PROTEIN. [1];	SCO-SPONDIN	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 79	Homo sepiens lysosomal-associated membrane protein 2 (LAMP2), transcript variant LAMP2A, mRNA	Homo sapiens lysosomal-associated membrane protein 2 (LAMP2), transcript variant LAMP2A, mRNA	Homo sapiens chromosome 21 segment HS21C004	D. melanogaster singed gene, exons 3, 4, 5 & 6	D. melanogaster singed gene, exons 3, 4, 5 & 6	C.perfringens plc gene for phosphalipase C upstream region containing bent DNA fragment	PROBABLE PEPTIDE ABC TRANSPORTER ATP-BINDING PROTEIN Y4TS	MR2-CT0222-211099-002-b10 CT0222 Homo sapiens cDNA	WR2-CT0222-211099-002-b10 CT0222 Homo sapiens cDNA	601676418F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3958997 5'	Arabidopsis thaliana mRNA for SigB, complete cds	Mus musculus T-cell receptor V region delta 1 chain gene, 5' region	Methanobacterium thermoautotrophicum from bases 702375 to 714311 (section 62 of 148) of the complete genome	Homo sapiens hHb5 gene for hair keratin, exons 1 to 9	Escherichia coli K-12 MG1655 section 225 of 400 of the complets genome	Mus musculus Emr1 mRNA, complete cds	Homo sapiens mysloid/lymphoid or mixed-lineage leukemia (trithorax (Drosophila) homolog); translocated to, 10 (AF10), mRNA	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 38	Mus musculus mennose receptor, C type 2 (Mrc2), mRNA	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 77	Homo sapiens GAP-like protein (LOC51306), mRNA	Homo sapiens GAP-like protein (LOC51306), mRNA	601811060R1 NIH_MGC_48 Horno sapiens cDNA clone IMAGE:4053851 3'
	Top Hit Database Source	LN LN	EST_HUMAN	EST HUMAN	SWISSPROT	N	LΝ	L	Ν	LN LN	۲	۲	SWISSPROT	EST_HUMAN	EST_HUMAN	EST_HUMAN	N	N	Į.	Z	۲	۲	F	Ľ.	ž	Z L	۲	Z	EST_HUMAN
9.6)	Top Hit Acession No.	Y10196.1	R94090.1	AW027174.1	P98167	AL161583.2	4504956 NT	4504956 NT	AL163204.2	X17550.1	X17550.1	X62825.1	Q53194	AW752901.1	AW752901.1	BE902390.1	AB004293.1	L41687.1	AE000856.1	Y19210.1	AE000335.1	U66888.1	11432598INT	AL161536.2	6678933 NT	AL161581.2	7706136 NT	7706136 NT	BF129796.1
	Most Similar (Top) Hit BLAST E Value		3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01	3.6E-01								3.6E-01	3.6E-01		3.5E-01	3.5E-01	3.5E-01	3.5E-01	
	Expression Signal	1.68	4.57	1.73	0.68	13.59	3.06	3.06	1.32	0.92	0.92	0.54	18.15	0.53	0.53	2.51	4.15	2.02	4.07	2.45	5.79	4.7	2.16	1.35	2.67	4.48	1.39	1.39	3.83
	ORF SEQ ID NO:	32008		32717	33623	33678	34388	34389	34597	34784	34785			35374	35375	36359	36531	36615	36025				:	25273	25369	25814			25935
	Exan SEQ 1D NO:		19733	19854	20707	20762	21470	21470	21656	21834	21834	21813				23344	23501	23577	23016	25109	24159	24258	24502	12791	12884	13327	13373		13430
	Probe SEQ ID NO:	6604	7202	7327	8166	8221	8932	8932	9120	9320	9320	0686	5225	9304	9904	10823	10987	11065	11318	11680	11768	11923	12308	120	223	208	754	754	812

Page 64 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Į	_	Т	Т	т —		Т	Т	Т	Т	T	T	1	T	T		T	Т	П	T	T	Т	П		Т	т	Т	1	Т	Т	т
	Top Hit Descriptor	Rattus norvegicus ADP-ribosylation factor-directed GTP ase activating protein mRNA, complete cds	HOMEOBOX PROTEIN HOX-A4 (HOX-1.4) (MH-3)	ZO8809.s1 Stratagene NT2 neuronal precursor 937230 Homo sapiens cDNA clone IMAGE:650872.3'	Fibrobacter succinogenes S85 endoglucanase E (celE) and endoglucanase D (celD) gene. complete cds	294f03.r1 Stratagene corneal stroma (#937222) Homo sapiens cDNA clone IMAGE:512285 5'	Info0d03 s1 NCI_CGAP_Lym3 Homo septens cDNA clone IMAGE:1172357 3'	Danio rerio homeobox protein (hoxb5b) gene, complete cds	788iE1 fetal brain cDNA Homo sapiens cDNA clone 788iE1-K similar to R07879, Z40498	Rat leukocyte common antigen (L-CA) gene, exons 1 through 5	EARLY E2A DNA-BINDING PROTEIN	EARLY E2A DNA-BINDING PROTEIN	Human mRNA for KIAA0086 gene, complete cds	PM4-SN0012-030400-001-a11 SN0012 Homo saplens cDNA	Zw79f03.r1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:782429 5' similar to TR:G1066935 G1086935 F10F2.1	Bos taurus peptide methionine sulfoxde reductase (msrA) mRNA, complete cds	GLUCOSE-6-PHOSPHATE 1-DEHYDROGENASE, CHLOROPLAST PRECURSOR (G6PD)	S scrofa mRNA for CD31 protein (PECAM-1)	Homo sapiens fumor protein p53-binding protein, 2 (TP53BP2), mRNA	RC4-ET0024-250600-014-d07 ET0024 Homo sapiens cDNA	Rattus norvegicus Na-K-Cl cotransporter (Nkcc1) mRNA, complete cds	Homo sapiens tyrosine kinase non-receceptor 1 (TNK1), mRNA	VOLTAGE-DEPENDENT N-TYPE CALCIUM CHANNEL ALPHA-1B SUBUNIT (CALCIUM CHANNEL, L TYPE, ALPHA-1 POLYPEPTIDE ISOFORM 5) (BRAIN CALCIUM CHANNEL III) (BIII)	X leevis gene for albumin including HP1 enhancer	QV2-HT0577-090400-128-c07 HT0577 Homo saplens cDNA	Cgriseus rhodopsin gene for opsin protein	Gallus gallus SPARC gene for ostbonectin, promoter and exon 1	Gallus gailus SPARC gene for ostaonectin, promoter and exon 1	yz90h12.r1 Soares_multiple_sclerosis_2NbHMSP Homo sapiens cDNA clone IMAGE:290375 5'	Human glucokinase (GCK) gene, repeat polymorphism
	Top Hit Database Source	Ę	SWISSPROT	EST_HUMAN	۲	EST HUMAN	EST_HUMAN	۲	EST_HUMAN	۲N	SWISSPROT	SWISSPROT	ΝŢ	EST_HUMAN	EST HUMAN	N L	SWISSPROT	Z	N.	EST_HUMAN	ΙN	¥	SWISSPROT	۲ <u>۷</u>	EST_HUMAN	۲	¥	N L	EST_HUMAN	LΝ
26	Top Hit Acession · No.	U35776.1	3.5E-01 P06798	01 AA223252.1	3.5E-01 U05897.1	3.5E-01 AA057691.1	3.5E-01 AA642138.1	01 AF071253.1	01 N81203.1	3.5E-01 M18349.1	096687	Q96687	3.5E-01 D42045.1	AW863916.1	3.5E-01 AA431833.1	3.5E-01 U37150.1	3.5E-01 O24357	X98505.1	11448042 NT	3.5E-01 BF358871.1	3.5E-01 AF051561.1	4507610 NT	002294	3.5E-01 Z26825.1	3.5E-01 BE174794.1	21 X61084.1	3.5E-01 AJ243178.1	3.5E-01 AJ243178.1	1 N77597.1	01 L05145.1
	Most Similar (Top) Hit BLAST E Value	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01 Q96687	3.5E-01 Q96687	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5€-01	3.55-01	3.5E-01	3.5E-01	3.5E-01 002294	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01	3.5E-01
	Expression Signal	1.91	1.12	1.92	11.8	0.57	1.27	2.3	0.57	4.33	0.74	0.74	1.42	6.0	0.75	0.72	0.93	3.51	2.02	0.85	0.61	1.12	1.93	4.91	1.14	4	2.09	2.09	1.93	1.71
	ORF SEQ ID NO:	26807	27469	27770		28124					28908	78508	31071		31932	31976	32172			33462		34319	35120	35284	35351	36157	38440			37050
	Exon SEQ ID NO:	14274	14893	15472	15284	15646	16476	16936	17588	17639	18169	18169	18364	18988	19139	19176	19363	19456	20558	20561	20943	21396	22150	22299	22374	23148	23422	23422	23912	23980
	Probe SEQ ID NO:	1682	2322	2637	2729	3030	3878	4349	5014	2066	5537	5537	5738	6384	6540	6578	6770	7116	8016	8019	8403	8857	9651	1086	248	10613	10902	10902	11462	11532

Page 65 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

			_	_		_	_		-	_	_	_	_	_	_				_			_	_	_	_			
Company of the compan	Top Hit Descriptor	Schistosoma mansori strain NMRI chromatin assembly factor 1 small subunit-ike protein (RBAP48) mRNA, complete cds	B. taurus atpA1 gene for F(0)F(1) ATP synthase alche-subunit	Thermologa maritima section 88 of 138 of the complete genome	Thermotoga maritima section 3 of 136 of the complete genome	ys84f11.r1 Soares retina N2b4HR Homo sapiens cDNA clone IMAGE:219597 5	ys64f11.r1 Soares retina N2b4HR Homo sapiens cDNA clone IMAGE:2195975	Homo sapiens partial N-myc (excn 3), HPV45 L2, HPV45 L1, HPV45 E6, HPV45 E7 and HPV45 E1 genes is dated from IC4 control continuous cell line.	Pseudomonas fluorescens colls colls agrees ad 222 and partial into American	QV3-HT0281-241199-019-010 HT0281 Home sapiens cDNA	Azotobacter vinelandii nifA gene for NifA protein (positive regulatory element)	Synechocystis sp. PCC8803 complete genome, 11/27, 1311235-1430418	Homo sapiens chromosome 21 segment HS21C010	Homo sapiens chromosome 21 segment HS21C010	Synechocystis sp. PCC6803 complete genome, 11/27, 1311235-1430418	Canis familiaris rod photoreceptor cGMP-cated channel alpha-subunit (CNGC1) mRNA commilers cuts	Homo sapiens pulmonary surfactant protein D. promoter region and exon 1	Methylovorus sp. strain SS1 putative GrpE (grpE), DnaK (dnaK), and putative DnaJ (dnaJ) genes, complete cds	7n94e01.x1 NCI_CGAP_Ov18 Homo sapiens cDNA clone IMAGE:3572232 3' similar to TR:Q9UJ15	Homo sapiens p47-phox (NCF1) gene, complete cds	no11b10.s1 NCI CGAP Phe1 Homo sapiens cDNA clone IMAGE:1100347 3'	Homo sapiens integrin alpha 6 (ITGA6) gene, exons 12 through 23	MR4-BT0403-230200-202-c01 BT0403 Homo sepiens cDNA	601901632F1 NIH_MGC_19 Homo saplens cDNA clone IMAGE:4130935 5'	qi95c05.x1 NCI_CGAP_Kid3 Homo sapiens cDNA clone IMAGE:1887208 3' similar to contains Alu repetitive element;	Arabidopsis thaliana DNA chromosome 4, conting fragment No. 90	zn12d11.s1 Stratagane hNT neuron (#837233) Homo sapiens cDNA clone IMAGE:5472213	Echovirus 22 1AB, 1C, 1D, 2A, 2B, 2C, 3A, 3B, 3C, 3D prateins RNA, complete mature peptides and cds
2001	Top Hit Database Source	Į	N	NT	Ľ.	EST_HUMAN	EST_HUMAN	Ę	L	EST HUMAN	ĽN	NŢ	NT	LN	NT	LN	N	۲N	FST HIMAN	TN	EST HUMAN	LN L	EST HUMAN	EST_HUMAN	EST HUMAN	Z	EST_HUMAN	NT
26	Top Hit Acession No.	AF297468.1	X64565.1	AE001774.1	AE001691.1	H80814.1	H80814.1	A.1242956 1	Y09798.2	AW380120.1	Y00554.1	D90909.1	AL163210.2	AL163210.2	D90909.1	U83905.1	AF034862.1	AF106835.1	BF449010 1	AF184614.1	AA584196.1	AF166341.1	BE069912.1	BF314689.1	AI240973.1	AL161594.2	AA085313.1	L02971.1
	Most Similar (Top) Hit BLAST E Value	3.5E-01	3.5E-01	3.5E-01				3.4F-01	_		3.4E-01	3.4E-01	3.4E-01	3.4E-01	3.4E-01		3.4E-01		3.45-01		3.4E-01/			3.4E-01	3.4E-01	3.4E-01	3.4E-01	3.4E-01
	Expression Signal	1.51	7.56	2.03	2.21	2.64	2.64	2	7.62	8.97	1.86	2.6	0.73	0.73	96.0	6.78	0.84	4.84	1 32	1.23	1.56	0.7	2	1.01	4.2	2.9	5.68	2.44
	ORF SEQ ID NO:						30626		28136			27584	!	28127	28268	28282	28473	28671						29795		31210		
	Exan SEQ ID NO:	25112	24211				24950	13356			13965				15796	15809	15996	16188	16456	16702	16718	17177	17330	17346	17660	18486	18599	18771
	Probe SEQ ID NO:	11778	11851	12014	12209	12643	12643	738	1011	1013	1371	2445	3032	3032	3183	3197	3387	3584	3858	4108	4126	4594	4749	4765	5087	5864	5979	6158

Page 66 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 67 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
12023	24865		3.18	3.4E-01	BE218652.1	EST_HUMAN	hv42h08.x1 NCI_CGAP_Lu24 Homo septens cDNA clone IMAGE:3176127 3' similar to contains PTR5.t3 PTR5 repetitive element;
12079	24974		2.44	3.4E-01	19838361 NT	N	Beta vulgaris mitochondrion, complete genome
12196	24424	30950		3.4E-01	AJ297131.1	L	Mus musculus SIL, MAP_17, CYP_a, SCL & CYP_b genes
12427	25068		1.25	3.4E-01	AJ288948.1	NT	Clostridium cellulolyticum partial spoIVB gene and spo0A gene, strain ATCC 35319
							Homo saplens HLA class III region containing tenascin X (tenascin-X) gene, partial cds; cytochrome P450 21-hydroxylase (CYP21B), complement component C4 (C4B) G11, halicase (SKI2W), RD, complement factor B
12523	24839		2.55	3.4E-01	AF019413.1	NT	(8f), and complement component C2 (C2) genes,>
12851	24723		2.71	3.4E-01	11466174 NT	NT	Naegleria gruberi mitochondrion, complate genome
18		25151	13.68	3.3E-01	X07990.1	IN	Rhizobium leguminosarum sym plasmid pRL5J1 nodX gene
110			3.75		X07990.1	IN	Rhizobium leguminosarum aym plesmid pRL5JI nodX gene
473					AL161545.2	INT	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 45
661	13285	25788	1.87	3.3E-01	7662485 NT	NT	Homo sapiens KIAA1100 protein (KIAA1100), mRNA
1242	13840	26358	2.96	3.3E-01	Q12446	SWISSPROT	PROLINE-RICH PROTEIN LAS17
1350	13945	26469	3.58		BF568880.1	EST_HUMAN	602184016T1 NIH_MGC_42 Homo sapiens cDNA clone IMAGE:4300251 3'
1649	14241	26775	1.43		6753685 NT	IN	Mus musculus disintegrin 5 (Dign5), mRNA
1773	14363		1.44	3.3E-01	AA332734.1	EST_HUMAN	EST36722 Embryo, 8 week I Homo capiens cDNA 5' end
2075	14855		1.22		AF031148.1	TN	Methylococcus capsulatus strain Bath outer membrane protein MopB (mopB) gene, complete cds
0370	15017		77 3		* 00E03*	, i	Homo saplens undine monophosphate synthetase (ordate phosphoribosy) transferase and ordidine-5-
35.55	1				*20/0C*	2	receiptoypasse (OWL 5) micked
9/82	15592	28074	2.14	3.3E-01	AJ251805.1	NT	Bacteriophage phi-YeO3-12 complets genome
3049	15865		99.0	3.3E-01	002743	SWISSPROT	INTERLEUKIN-12 ALPHA CHAIN PRECURSOR (IL-12A) (CYTOTOXIC LYMPHOCYTE MATURATION FACTOR 35 KD SUBUNIT) (CLMF P35)
3091	15708	28178	0.82	3.3E-01	AJ007932.2	IN	Streptomyces argillaceus mithramycin biosynthetic genes
3542	16147	28629	66'0	3.3E-01	AB012922.1	NT	Homo saplens MTA1-L1 gene, complete cds
3882	16480	28942	2.14	3.3E-01	084645	SWISSPROT	EXODEOXYRIBONUCLEASE V BETA CHAIN
							GENOME POLYPROTEIN (CONTAINS: N-TERMINAL PROTEIN (P1); HELPER COMPONENT
3890				3.3E-01	P22602	SWISSPROT	PROTEINASE (HC-PRO); PROTEIN P3]
4037				3.3E-01	AL161498.2	LN	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 10
4073	16669	29130	1.95	3.3E-01	AF200448.1	NT	Нурохуюл fregiforme chitin synthase gene, partial cds
4457	17043		1.44	3.3E-01	D31662.1	NT	Rattus norvegicus DNA for regucalcin, partial cds
4799	17377		1.57	3.3E-01	AI539114.1	EST HUMAN	tp78b12.x1 NCI_CGAP_UI3 Homo sapiens cDNA clone IMAGE:2205407 3' similer to gb:X57522 ANTIGEN PEPTIDE TRANSPORTER 1 (HUMAN);
4964	1	29980		3.3E-01	D64003.1	NT	Synachocystis sp. PCC8803 complete genome, 22/27, 2755703-2868766
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Page 68 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Page 69 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 61	Fuserium pose virus 1 RNA2 putative RNA dependent RNA polymerase gene, complete cds	P.vulgaris arc5-1 gene	I LACTOSE PERMEASE (LACTOSE-PROTON SYMPORT) (LACTOSE TRANSPORT PROTEIN)	S.cerewisiee chromosome II reading frame ORF YBR172c			Botry dis cinerea strain T4 cDNA library under conditions of nitrogen deprivation	4 601868804F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:41115125	Mus musculus Pbxknotted 1 homeobox (Pknox1), mRNA	Homo sapiens promyelocytic leukemia zinc finger protein (PLZF) gene, complete cds	Humam h NAT allele 3-2 gene for anytamine N-acetytransferase	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 46	Rabbit beta-like globin gene cluster encoding the epsilon, gamma, delta (pseudogene) and beta globin			Г		Т	Homo sapiens interleukin 12 p40 subunit (IL12B) gene, IL12B-1 allele, complete cds	Ī	Giardia intestinalis pyruvate flavodoxin oxidoreductase and flanking genes	Human mRNA for KIAA0361 gene, KIAA0361 protein	Rat ISO-atrial natriuretic factor gene, complete cds	Rattus norwegicus repeat, map NOS-D12W ox1	H.sapiens gene fragment for acety/choline receptor (AChR) alpha subunit exons 8.9 and 3' flanking region	Т	Arabidopsis thaliana DNA chromosome 4, contig fregment No. 70	1 801855580F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4075627 5'		Deinococcus radiodurans R1 section 152 of 229 of the complete chromosome 1	Oryctolagus, cuniculus ig H-chain pseudogene, V-region (VH6-a2) gene, partial cds
Top Hit Detabase Source	Ž	Σ	ΝŢ	SWISSPROT	Z L	EST_HUMAN	EST_HUMAN	¥	EST_HUMAN	ΓN	NT	N N	N		NT	SWISSPROT	EST_HUMAN	SWISSPROT	EST HUMAN	F	EST_HUMAN	IN	IN	IN	ΙN	Ę	EST HUMAN	Ā	EST_HUMAN	EST_HUMAN	IN	ΙN
Top Hit Acession No.	AL161561.2	AF047013.1	250202.1	Q48624	236041.1	AW957194.1	AW957194.1	AL111655.1	BF203817.1	TN 6700177	AF060568.1	D10872.1	AL161546.2		M18818.1	010268	BF693617.1	057081	BE782748.1	AY008847.1	BE173964.1	1.127221.1	AB002359.1	M60266.1	AJ231001.1	X02508.1	BF311635.1	AL161574.2	BF246771.1	BF246771.1	AE002015.1	U51026.1
Most Similar (Top) Hit BLAST E Value	3.2E-01	3.2E-01	3.2E-01	3.2E-01		3.2E-01	3.2E-01	3.2E-01	3.2E-01	3.2€-01		3.2E-01	3.2E-01				3.2E-01	3.2E-01		3.2E-01	3.2E-01			3.2€-01	3.2E-01	3.2E-01	3.2E-01	3.2E-01	3.2€-01	3.2€-01	3.2E-01	3.2E-01
Expression Signal	0.78	27.98	1.39	7:37	0.92	6.36	6.36	1.22	2.89	2.24	1.09	0.78	0.61		1.64	1.56	6.52	0.63	0.58	0.93	2.5	1.18	1.03	1.33	0.51	11.34	16.78	1.43	1.22	1.22	2.72	0.69
ORF SEQ ID NO:		26317	28441	26555	26946				27354		27857					29628		30025			30519	31481		33563	33665	33772	33777					34058
Exan SEQ ID NO:	13367	13804	13919	14027	14401	LJ		14469				16269	16621				17422	17582	Ι,	17950			19421		20751	20849	20852	20938				21144
Probe SEQ ID NO:	747	1204	1325	1434	1811	1819	1819	1883	2205	2578	2734	3668	4023		4483	4597	4844	2009	5174	5392	5478	6112	6831	8113	8210	8308	8311	8388	8437	8437	8508	8605

Page 70 of 526 Table 4

Homo sapiens 8-phosphofructo-2-kinase/fructose-2,6-bisphosphatase (PF2K) gene, exons 12 and 13 Homo sapiens 6-phosphofructo-2-kinasie/fructose-2,6-bisphosphatase (PF2K) gene, exons 12 and 13 re90h06.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:125051 5' similar to Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds; and unknown genes EST04702 Fetal brain, Stratagene (cat#936206) Homo sapiens cDNA clone HFBDZ21 hi46h08.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2975391 3 Oryctolagus cuniculus lg H-chain pseudogene, V-region (VH6-a2) gene, partial cds H.sapiens gene for immunoglobulin kappa light chain variable region A8 and A9 MR2-CT0222-281099-005-h05 CT0222 Homo saplens cDNA hv89f05.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE:3181569 3 601275480F1 NIH_MGC_20 Homo sapiens cDNA clone IMAGE:3616746 5 ql39d01.x1 NCI_CGAP_Co8 Hamo sapiens cDNA clone IMAGE:1874689 3 Bos taurus inositol 1,4,5-trisphosphate receptor type I mRNA, complete cds Homo sapiens hepatocyte nuclear factor-3 alpha (HNF3A) gene, exon 1 carboric anhydrase IV [rats, Sprague-Dawley, lung, mRNA, 1205 nt] Drosophila melanogaster laminin A (Lam-A) mRNA, complete cds **Top Hit Descriptor** Arabidopsis thaliana DNA chromosome 4, contig fragment No. Human monocyte antigen CD14 (CD14) mRNA, complete cds RC3-HN0001-310300-011-b04 HN0001 Homo saplens cDNA Homo sapiens deoxycytidylate deaminase gene, complete cds Daucus carota mRNA for transcription factor E2F (E2F gene) Xytella fastidiosa, section 130 of 229 of the complete genome Homo sepiens KiAA0174 gene product (KIAA0174), mRNA Homo sapiens KIAA0174 gene product (KIAA0174), mRNA S.cerevisiae chromosome XV reading frame ORF YOL141w Homo sapiens filamin 2 (FLN2) gene, exons 10 through 22 Mus musculus gene for Ser/Thr kinase KKIAMRE, exon 6 Homo sapiens chromosome 21 segment HS2 Homo sapiens gene for AF-6, complete cd: ELONGATION FACTOR TU (EF-TU) gb:M64241 QM PROTEIN (HUMAN); Mus musculus mRNA for polycystin Single Exon Probes Expressed in Fetal Liver EST_HUMAN Top Hit Database EST_HUMAN EST HUMAN EST HUMAN EST HUMAN EST_HUMAN HUMAN EST HUMAN Source EST Ę ¥ ż 뉟 ż Top Hit Acession 7661971 7661971 3.1E-01 AW629036.1 3.1E-01 AW377354.1 3.2E-01 AL163204.2 3.2E-01 M86511.1 3.1E-01 AE003984.1 3.1E-01 AW 983549.1 3.2E-01 BE326230.1 3.1E-01 AF176111.1 3.2E-01 AF041829.1 3.2E-01 AF041829.1 3.2E-01 BE385776.1 AL161503.2 3.1E-01 AF184122.1 3.2E-01 AB011399. 3.1E-01 AB029069. AI264458.1 ģ AJ251586. 3.1E-01 S68245.1 R18051.1 Z74883.1 3.2E-01 U51026.1 J44914. 3.2E-01 L07288.1 3.1E-01|X71887.1 3.2E-01|L39874. 3.2E-01 083217 3.2E-01 3.1E-01 3.1E-01 3.1E-01 3.1E-01 3.1E-01 3.1E-01 3.1E-01 3.1E-01 (Top) Hit BLAST E 0 8 89 0.58 0.48 3.03 3.28 4.31 3.37 8 8 2.39 3.53 0.79 0.82 96'0 10.8 0.75 0.99 263 2.08 0.51 3.67 3.67 0.91 9.0 Expression Signal 34465 34548 35588 27823 27843 30072 30109 30194 31295 31892 32052 34059 34547 35376 29046 32191 ORF SEQ 36098 30504 31127 Ö Q 21144 21535 21613 21613 24999 24525 24819 24618 15820 17628 17670 18296 18411 18565 19189 19375 25041 15403 16578 19597 SEQ ID 22402 15403 15501 17771 Ę ģ 10548 11796 12344 2883 3208 6784 6863 8605 2077 9905 10100 10213 12489 12550 5055 5087 5206 5669 5786 5945 SEO ID 8997 2695 2722 2722 507 6592 6654 80/2 ö

Page 71 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Probe SEQ ID NO: 7048 7671 8882 9979 9979 11024 11249 12535 12635 12646	^ω σ	g, Ö	Signess	Mosi Mosi (1)	Top Hit Ad No No No No No No No No No No No No No	THUMAN THUMAN THUMAN THUMAN THUMAN THUMAN THUMAN THUMAN	Top Hit Describtor Challesse Scurce Source S
5695 5695			4.03	3.0E-01 3.0E-01	01 BE693575.1 01 BE693575.1	EST_HUMAN EST_HUMAN	RC3-BT0333-190700-111-e03 BT0333 Home sapiens cDNA RC3-BT0333-190700-111-e03 BT0333 Home sapiens cDNA
5731	1 1	Ш	4.57	3.0E-01		N 1	Mus musculus 129/sv Clara cell 10 kd protein (mCC10) gene, complete cds Morse cyddagain 15 gans complete cds
20			٥. ١٣١	3.00.0			Mouse cytokeratin 15 gene, complete cds

Page 72 of 526

p21a11.x1 NCI_CGAP_Gas4 Homo sapiens cDNA clone IMAGE:2188412 3' similar to gb:D15050 NIL-2-A Mus musculus C-type (calcium dependent, carbohydrate recognition domain) lectin, superfamily member 9 2557d12.r1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:701591 5' similar to contains Alu Anabaena PCC7120 cytosine-specific DNA methyltransferase (dmnB) gene, complete cds; putative Rattus norvegicus mRNA for glyceraldehyde-3-phosphate dehydrogenase type 2 (gapdh-2 gene) yp84b10.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:194107 5 yp84b10.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:194107 5' Homo sapiens membrane component, chromosome 11, surface marker 1 (M11S1) mRNA xe03d10.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2606035 3 Strongylocentrotus purpuratus 34/67 kDa laminin-binding protein mRNA, partial cds HYPOTHETICAL 59.5 KD PROTEIN IN WZA ASMA INTERGENIC REGION 602133271F1 NIH_MGC_81 Hamo sapiens cDNA clone IMAGE:4288336 5 Streptomyces sulfondaciens isopenicillin N synthase (pcbC) gene, partial cds wr02f10.x1 NCI_CGAP_GC8 Homo sepiens cDNA clone IMAGE:2480395 3 602140133F1 NIH_MGC_46 Homo sapiens cDNA clone IMAGE: 4301097 5 601339079F1 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3681594 5' 602140133F1 NIH_MGC_48 Homo sapiens cDNA clone IMAGE:4301097 5' anthranilate phosphoribosytransferase gene, partial cds; and unknown gene ZINC FINGER PROTEIN (HUMAN) contains element L1 repetitive element Streptococcus pneumoniae strain DBL5 PspA (pspA) gene, partial ods Homo sapiens DKFZP586M0122 protein (DKFZP586M0122), mRNA Aspergillus oryzae bipA gene for ER chaperone BiP, complete cds Thermotoga maritima section 67 of 136 of the complete genome Top Hit Descriptor Mus musculus ribose 5-phosphate Isomerase A (Rpia), mRNA Mouse apolipoprotein A-II (Alp.2) gene, complete cds PM1-CT0326-171299-001-f12 CT0326 Homo sapiens cDNA PM1-CT0326-171299-001-f12 CT0326 Homo sepiens cDNA Aquifex aeolicus section 68 of 109 of the complete genome Cantagalo orthopoxvirus hemagglutinin gene, complete cd. Homo sapiens chromosome 21 segment HS21C006 Mus musculus midnolin (Midn-pending), mRNA Single Exon Probes Expressed in Fetal Liver PONTICULIN PRECURSOR epetitive element Clecsf9), mRNA EST_HUMAN EST_HUMAN EST_HUMAN EST_HUMAN HUMAN EST HUMAN EST HUMAN EST_HUMAN HUMAN EST HUMAN HUMAN EST_HUMAN SWISSPROT Top Hit Database SWISSPROT Source EST EST EST_ F Ę 눋 뉟 눋 Ż 뉟뉟 9910161 NT 6677766 NT 5174502 NT 10947007 7661685 Top Hit Acession 3.0E-01 AW118111.1 AW 754239.1 AW 754239.1 AW002902.1 BE566083.1 2.9E-01 AA284468.1 3.0E-01 AF141676.1 AF220507.1 BF574612.1 3.0E-01 AL163206.2 3.0E-01 AF071810.1 BF683841.1 BF683841.1 AE001755.1 AI610836.1 3.0E-01 AF229247.1 ĝ 2.9E-01 M32360.1 H51029.1 H51029. 3.0E-01 P54660 3.0E-01 3.0E-01 3.0E-01 3.0E-01 2.9E-01 2.9E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 3.0E-01 2.9E-01 3.0E-01 (Top) Hit BLAST E 3.0E-0 0.73 1.38 1.21 96.0 0.82 0.98 8 0.56 1.95 0.73 1.37 251 2.07 0.71 8 6 Expression Signal 28379 29609 30475 32322 32555 32756 35778 35802 29032 32899 33316 33870 34223 34624 35780 35801 27064 ORF SEQ Ö N O 17166 16563 21680 15900 SEQ ID 18052 19503 19707 19893 20409 20855 20951 21302 21344 22790 22809 24047 24047 24287 24984 25033 14509 15088 16751 20033

8314

7867

8763 8805

SEQ ID

7005 7175 7512

6944

9145 9493 9878

10315

11604 11604 11975 2529

10315

10296

4583

3965 4159

3289

1924 2070

3

Page 73 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

	_								-		_	_	-	_	_	_	_	_	_		_	-	_		_	_		_		_		۰.,
Top Hit Descriptor	Hamo sapiens chramosome 21 sogment HS21C007	Lymantia dispar vitellogenin gene, complete cds	Homo sapiens KIAA0537 gene product (KIAA0537), mRNA	we06f03.x1 NCI_CGAP_Kid11 Homo sepiens cDNA clone IMAGE:2297309 3' similar to contains L1.t2 L1	repetitive element;	y77e12.s1 Soares infant brain 1NIB Homo saplens cDNA clone IMAGE:28291 3'	Suaeda maritima subsp. salsa S-adenosylmethionine sythetase 2 mRNA, complete cds	B. subtilis levenase operon levD, levE, levC and secC (partial) genes for fructose phosphotransferase	System polypopudos F 10, 10, 20, 30 and 10 years a	B.subblis levanase operon levD, levE, levF, levG and sacC (partial) genes for fructose phosphotransferase system polypeptides P16,18,28,30 and levanase	Mus muscutus Eph receptor A8 (Epha8), mRNA	zv97b12.r1 Soares_NhHMPu_S1 Homo sapiens cDNA clone IMAGE:767711 5'	we27c05.x1 NCI_CGAP_Lu24 Homo sepiens cDNA clone IMAGE:2342312 3' similar to contains L1.t1 L1	repetitive element;	Bos faurus myosin I mRNA, complete cds	y/39d08.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:141915 5'	y/39d08.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:141615 5'	Mus musculus Flith protein (Flith) gene, complete cds; and Light protein (Light) gene, partial cds	PUTATIVE MULTICOPPER OXIDASE YDR506C	Mus musculus major histocompatibility locus class II region; Fas-binding protein Daxx (DAXX) gene, partial cds; Bing1 (BING1), tapasin (tapasin), RalGDS-like factor (RLF), KE2 (KE2), BING4 (BING4), beta1, 3-	galactosyl transferase (beta1,3-galactosyl tr>	601065830F1 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3452287 5'	601065830F1 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3452287 5'	801882570F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4095113 5	AU150910 NT2RP2 Homo sepiens cDNA clone NT2RP2003901 3'	Arabidopsis thaliana sulfonylurea receptor-like protein mRNA, complete cds	Baboon lymphocyte homing/adheslon receptor mRNA, complete cds	Pyrococcus abyasi complete genome; segment 5/6	Pyrococcus abyasi complete genome; segment 5/6	Trypanosoma cruzi stage-specific surface glycoprotein gp82 (gp82) mRNA, partial cds	Torpado californica mRNA encoding acetylcholine receptor gamma subunit	Torpado californica mRNA encoding acetylcholine receptor gamma subunit
Top Hit Database Source	NT	IN	۲		EST_HUMAN	EST_HUMAN	LN.	. H	2	F	FZ	EST_HUMAN		EST_HUMAN	IN	EST_HUMAN	EST_HUMAN	IN	SWISSPROT		F	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN	LN	Ņ	LN	LN	NT	TN	NŢ
Top Hit Acession No.	2.9E-01 AL163207.2	J90756.1	7662169 NT		AI670899.1	2.9E-01 R37485.1	4F321001.1	2 0E 04 VE6008 4	730030 I	2.9E-01 X56098.1	6879682 NT	2.9E-01 AA418145.1		٠,		2.9E-01 R69194.1		1.63	204399		2.9E-01 AF100956.1	2.9E-01 BE540422.1	2.9E-01 BE540422.1	2.9E-01 BF217743.1	2.9E-01 AU150910.1	2.9E-01 AF225908.1	M22452.1	2.9E-01 AJ248287.1	2.9E-01 AJ248287.1	2.9E-01 AF128843.1	2.9E-01 V01394.1	2.9E-01 V01394.1
Most Similar (Top) Hit BLAST E Value	2.9E-01	2.9E-01	2.9E-01		2.9E-01	2.9E-01	2.9E-01	2000	4.0E-01	2.9E-01	2.9E-01	2.9E-01		2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01 Q04399		2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01
Expression Signal	0.63	1.02	1.43		1.7	1.25	0.79	ű	5	5.1	90'9	1.26		0.93	2.3	0.72	0.72	1.35	2.87		1.61	1.92	1.92	68.0	99.0	1.07	0.71	77.0	72.0	2.24	2.88	2.88
ORF SEQ ID NO:			30155				32344	900.70		31287	31302						31984	30437	32527		32598		33309	L	33990			34689	34690	36302		36590
Exon SEQ ID NO:	17372	17718	17724		17847	18098	19522	10550	200	18558	18570	18816		19030	19068	19183	19183	18081	19885		19744	20402	20402		i	21395		21747			23555	23555
Probe SEQ ID NO:	4793	5148	5154		5285	5463	5592	6097	200	5937	5949	. 6206		6427	6487	6585	6585	7062	7153		7213	7860	7860	8103	8531	8856	8963	9170	9170	10773	11041	11041

Page 74 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	ny35h02.s1 NCI_CGAP_Pr12 Homo sapiens cDNA clone IMACE:1273779 similar to contains LTR8.t2 LTR8 AN repetitive element;	Campylobacter jejuni NCTC11168 complete genome; segment 5/8		T	Г		Chlamydomonas reinhardtii mRNA for nitrite reductase structural tocus	Rattus norvegicus A-kinase anchoring pratein AKAP150 mRNA, complete cds	Prune dwarf virus movement protein, complete cds; coat protein, complete cds	Guira guira cocyte maturation factor Mos (c-mos) gene, partial cds	Г	П		AN QV1-CT0364-120200-065-b05 CT0364 Home sapiens cDNA	AN DKFZp586i2321_r1 586 (synonym: hute1) Homo sapiens cDNA clone DKFZp586i2321	AN hd44b03.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2912333.3	Escherichia coli K-12 MG1655 section 384 of 400 of the complete genome	Escherichia coli K-12 MG1655 section 384 of 400 of the complete genome	Arabidopsis thaliana DNA ctromosome 4, contig fragment No. 65	Arabidopsis thaliana mRNA for lipoyltransferase, complete cds	Toxoplasma gondii 90kDa heat-shock protein (HSP90) mRNA, pertial cds	B.taurus microsatellite (ETH121)	B.taurus microsatellite (ETH121)	Pyrococcus horikoshii OT3 genomic DNA, 777001-994000 nt position (4/7)	Borrelia burgdorferi (section 66 of 70) of the complete genome	Pseudomonas aeruginosa PA01, section 11 of 529 of the complete genome	ov44g10.x1 Scares_testis_NHT Homo sapiens cDNA clone IMAGE:1640226 3' similar to contains Alu	AN repetitive element; contains element MER22 repetitive element;	Mus musculus chromosome X contigA; putative Mages9 gene, Caltractin, NAD(P) steroid dehydrogenase	and Zinc tinger protein 185 OT IDNA DOLIVAKEDASE BETA SI IBI INIT // ABCE STOLICTI IDAL DOCTEIN // DOCTEIN /	Т	Human mKNA for transcription factor AREB6, complete cds
Top Hit Database Source	EST_HUMAN	ΙN	EST HUMAN	N	EST_HUMAN	ΙN	١	ΙN	١	<u></u> L	EST_HUMAN	EST HUMAN	۲	EST_HUMAN	EST_HUMAN	EST_HUMAN	IN	N L	ΙN	NT	N	IN	IN	LN	F	۲		EST_HUMAN		NI	SVISSIVE.	ź
Top Hit Acession No.	1 AA935373.1	1 AL 139078.2	AW005671.1	AF092453.1	2.9E-01 BE788199.1	2.9E-01 Y08937.1	1 Y08937.1	U67136.1	L28145.1	AF168050.1	BE313442.1	BE313442.1	D86550.1	AW 860020.1	AL047620.1	AW511195.1	AE000494.1	AE000494.1	AL161565.2	AB020975.1	AF179480.1	Z14037.1	Z14037.1	AP000004.1	AE001180.1	AE004450.1		AI090868.1		AL021127.2	210217	2.8E-01 D15050.1
Most Similer (Top) Hit BLAST E Value	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.9E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01	2.8E-01		2.8E-01		2.8E-01	2.05-01	2.8E-U1
Expression	2.07	5.52	1.54	1.47	4.4	1.57	1.57	2.06	0.75	3.14	3.51	3.51	1.03	2.01	2.12	3.53	2.41	2.41	2.75	1.21	1.7	2.36	2.36	1.26	2.06	0.62		2.75	,	1.32	12.7	1.07
ORF SEQ ID NO:		36973	30944			30877	30878			26238		26436	26448		27210	27322		27649		27813			28095	28513	29125					20500	l	28828
Exan SEQ ID NO:		23906	24411	24472	24505	24679	24679	13224	13228	13725	13914	13914	13928	14355	14638	14752	15075	15075	15147	15246	15614	15615	15615	16033	16664	16791		16862		17142		1/4/2
Probe SEQ ID NO:	11453	11456	12172	12262	12313	12586	12586	594	299	1122	1320	1320	1334	1765	2027	2175	2511	2511	2584	2688	2888	2999	2999	3425	4068	4202		4276	,	4559	1995	4897

Page 75 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 76 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Fujinami sarcona virus, complete genome	601654822R1 NIH_MGC_57 Hamo sapiens cDNA clane IMAGE:3839765 3'	601880794F1 NIH_MGC_55 Homo sapiens cDNA clone IMAGE:4109350 5	601880794F1 NIH_MGC_55 Homo sepiens cDNA clane IMAGE:4109350 5'	601852148F1 NIH_MGC_56 Homo sepiens cDNA clone IMAGE:4076028 5'	Drosophila heteroneura fruitless (fru) gene, alternative splice products, 5' flanking region, exons 1 through 7	and complete cds	602137418F1 NIH_MGC_83 Homo sapiens cDNA clone IMAGE:4273853 5'	Mus musculus DNA for prostaglandin D2 synthase, complete cds	PM4-HT0606-030400-001-s07 HT0606 Homo sapiens cDNA	601673020F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3955996 5'	Homo sapiens CDC42-binding protein kinase beta (DMPK-lika) (CDC42BPB), mRNA	Rattus norvegicus CDK104 mRNA	zx39b10.s1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone IMAGE:788827 3' similar to	contains Alu repetitive element,	Ipomoea purpurea transposable element Tip100 gene for transposase, complete cds	G.lamblia SR2 gene	2d22h10.r1 Soares_fetal_heart_NbHH19W Homo sapiens cDNA clone IMAGE:341443 5'	GAG POLYPROTEIN (CONTAINS: INNER COAT PROTEIN P12; CORE PROTEIN P15; CORE SHELL PROTEIN P30; NUCLEOPROTEIN P10]	Raffus norregicus vesicular monoamine transporter type 2, promoter region and exon 1	Feline immunodeficiency virus env gene, isolate ITTO088PIU (M88), partial	te43c11.x2 NCI_CGAP_Lu25 Homo sepiens cDNA clone IMAGE:2046838 3' simitar to contains element L1	repetitive element;	CM1-HT0875-060900-385-e05 HT0875 Homo sepiens cDNA	wc92e11.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2482828 3'	Drosophila buzzati alpha-esterase 6 (aE6) gene, partial cds	Drosophila buzzatii alpha-esterase 6 (aE6) gene, partial cds	Homo saplens DiGeorge syndrome critical region, telomeric end	Triticum aestivum (Wcs66) gene, complete cds	RC1-CT0286-230200-016-e03 CT0286 Homo sepiens cDNA	HOMEOBOX PROTEIN HOX-A4 (CHOX-1.4)	Astreopora myriophthalma mitochondrial cytb gene for cytochrome b, partial cds	Archaeoglobus fulgidus section 13 of 172 of the complete genome
Top Hit Database Source	NT	EST_HUMAN	EST_HUMAN	EST_HUMAN	EST_HUMAN			EST HUMAN	IN	EST_HUMAN	EST_HUMAN	NT	NT		EST_HUMAN	IN	IN	EST_HUMAN	SWISSPROT	NT	LN FN		EST_HUMAN	EST_HUMAN	EST_HUMAN	IN	. IN	NT	IN	EST_HUMAN	SWISSPROT	NT	TN
Top Hit Acession No.	9626154 NT	BE959727.2	BF241062.1	BF241062.1	BF695970.1		AF051662.1	BF674023.1	D83329.1	BE178699.1	BE900116.1	11433629 NT	Y17324.1		AA450061.1	AB004906.1	X79815.1	W58067.1	P03341	AF047575.1	Y13868.1		AI310858.1	BF088284.1	AI928015.1	AF216214.1	AF216214.1	L77569.1	L27516.1	AW856131.1	P17277	AB033171.1	AE001094.1
Most Similar (Top) Hit BLAST E Value	2.8E-01	2.8E-01			2.8E-01		=1					2.8E-01	2.7E-01		ĺ	2.7E-01		2.7E-01		2.7E-01	2.7E-01			į		2.7E-01	2.7E-01	-	2.7E-01	2.7E-01	2.7E-01		2.7E-01
Expression Signal	0.81	0.47	2.26	2.26	2.83		3.31	4.56	15.74	8.89	1.25	2.21	3.21		2.53	1.69	2.17	3.34	4.14	2.77	7.35		3.82	0.73	1.98	62.0	0.79	2.31	0.98	3.82	2.07	96'0	1.07
ORF SEQ ID NO:		35738	36166	36167	36197		36297				30927		25622		25740	26418		26903	26948		27545		27634		29138	29147	29148	29151	30037		30424		32122
Exon SEQ ID NO:	22710	22750			23183		23284					24988	13134			13898		14357	14403	15459	14973			Ⅎ			1						19318
Probe SEQ ID NO:	10215	10255	10622	10622	10651		10760	11158	12213	12328	12356	12519	502		ङ	1304	1662	1767	1813	2181	2405		2496	3013	4082	4096	4096	4101	2020	5193	5471	5681	6724

Page 77 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Archaeoglobus fulgidus section 13 of 172 of the complete genome FIBRILI IN 1 PRECIRSOR	Drosophila melanogaster rfc40 protein, Rop protein (Rop), and small GTP binding protein (DRas2) genes, complete cds.	HYPOTHETICAL 20.9 KD PROTEIN B0563.3 IN CHROMOSOME X	NITROGEN REGULATORY PROTEIN NUT1	NITROGEN REGULATORY PROTEIN NUT1	Bos taurus micromolar calcium activated neutral protease 1 (CAPN1) gene, exons 11-20, and partial cds	Bos taurus micromolar calcium activated neutral protease 1 (CAPN1) gene, exons 11-20, and partial cds	EST58740 Infant brain Homo sepiens cDNA 5' end similar to similar to myxsin-binding protein H	EST58740 Infant brain Homo saplens cDNA 5' end similar to similar to myosin-binding protein H	ze35b11.s1 Socres retina N2b4HR Homo septens cDNA clone IMAGE:380957 3' similer to contains Alu repetitive element;	Carasslus auratus pituliary adenylate cyclase activating polypeptide type 1 receptor precursor mRNA, complete cds	MR1-SN0062-100500-002-d09 SN0062 Homo sapiens cDNA	yc91h06.s1 Soares infant brain 1NIB Homo saplens cDNA clone IMAGE:23511.37	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 52	MAJOR VAULT PROTEIN (MVP) (LUNG RESISTANCE-RELATED PROTEIN)	Staphylococous aureus transposon Tn554	THREONYL-TRNA SYNTHETASE (THREONINE-TRNA LIGASE) (THRRS)	THREONYL-TRNA SYNTHETASE (THREONINE-TRNA LIGASE) (THRRS)	FIMBRIAE W PROTEIN	Rattus norvegicus DNA for peroxisome assembly factor-2, exon 4, 5, 6, 7, 8, 9, 10, 11,12, 13, 14, 15, 16, 17 and complete cds	Oryctolagus cuniculus calgranulin C mRNA, partial cds	Mus musculus transcription factor NF-ATc isoform a (NF-ATca) mRNA, complete cds	Homo sapiens xeroderma pigmentosum complementation group C (XPC) gene, intron 9	Homo sapiens xeroderma pigmentosum complementation group C (XPC) gene, intron 9	AV705043 ADB Hamo sapiens cDNA clone ADBCOD05 5'	AV705043 ADB Homo seplens cDNA clone ADBCOD05 5'
Top Hit Database Source	NT	LZ	SWISSPROT	SWISSPROT	SWISSPROT	NT	LN	EST HUMAN		EST_HUMAN	Ę	EST HUMAN		L	SWISSPROT		SWISSPROT	SWISSPROT	SWISSPROT	. LN	ΓN	FZ				EST_HUMAN
Top Hit Acession No.	AE001094.1 O81554	U15967.1	Q11079	001168	Q01168	AF248054.1	AF248054.1	AA351121.1	AA351121.1	AA013147.1	AF048820.1	AW 868503.1	R39257.1	AL161552.2	Q14784	X03218.1	083809	083809	P37928	D89660.1	AF091848.1	AF087434.1	AF156539.1	AF156539.1	AV705043.1	AV705043.1
Most Similar (Top) Hit BLAST E Value	2.7E-01	2.7E-01			2.7E-01	2.7E-01	2.7E-01	2.7E-01	2.7E-01		2.7E-01		_	2.7E-01	2.7E-01	2.7E-01	2.7E-01	2.7E-01	2.7E-01	2.7E-01	2.7E-01		2.7E-01		2.7E-01	2.7E-01
Expression Signal	1.07	0.78	0.87	0.95	0.95	2.21	2.21	0.92	0.92	0.95	0.51	0.59	0.48	0.94	0.83	0.53	8.93	9.93	2.02	0.67	16.0	2.5	0.69	0.69	2.31	2.31
ORF SEQ ID NO:	32123		32782		32955	33076	33077	33116		33258	-	33534	33577	33694				34735		35188	35476	35517	35834	35635	38233	36234
Exen SEQ ID NO:	19318	l		20079	2002	20188	20188	Ι.	20228	20348	20511	20821			21233				21785	22214	22488	22521	22643	22643	23221	23221
Probe SEQ (D NO:	6724 6875	7085	7383	7582	7562	7877	7677	7720	7720	7805	7969	6208	8127	8232	8694	8957	9258	9228	9228	9718	8663	10028	10148	10148	10691	10891

Page 78 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Exon SEQ ID NO: 23231 24863 24627 15416 114078 14078 14773 16243 16243 17034 17034 17034 17034	ORF SEQ ID NO: 28515 28616 27080 27081 28719 28775 29274 29622 29684 29684 29684 29684 29684 29684 29684	Signel Si	Most Similar (Top) Hit BLAST E Vatue 2.7E-01 2.7E-01 2.6E-01		Top Hit Database Source NT NT SWISSPROT NT SWISSPROT NT EST HUMAN NT NT NT NT NT NT NT NT NT NT NT NT NT N	Homo sapiens caveolin-1/-2 locus, Contig1, D75522, genes CAV2 (exons 1, 2a, and 2b), CAV1 (exons 1 and 2b) 2) 24 25 26 27 28 28 29 29 29 29 29 29 29 29
4752 17333 4825 17403	29776	1.63		U01103.1 AF142703.1	TN TN	Arabidopsis thalisma PSI type III chlorophyll a/b-binding protein (Lhca3*1) mRNA, complete cds Ophrestia radicosa maturase-like protein (matt\) gene, complete cds; chloroplast gene for chloroplast product
5107 17679 5195 17760 5544 18176 5640 18269	30742	3.56 0.58 1.29 0.68		H04858.1 AA884625.1 AB035972.1 M96060.1	EST_HUMAN NT NT	y51e05.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:152288 5 am33b11.s1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:1488605 3' Paramecium caudatum gene for PAP, complete cds Acetobacter xylinum collulose synthase (bcsA) gene, partial cds, CMCax and CcpAx genes, complete cds

Page 79 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	ta16a03.x1 NCI_CGAP_C018 Homo saptens cDNA clone IMAGE:2075788 3' similar to contains element MER35 repetitive element;	Homo sapiens protein translocase, JM26 protein, UDP-galactose translocator, pim-2 protoncogene homolog pim-2h, and shal-type potassium channel genes, complete cds; JM12 protein and transcription factor IGHM enhancer 3 genes, partial cds; and unknown g>	Thermotoga maritima section 123 of 138 of the complete genome	ts02e12.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2227438 3' similar to SW:NDF1_RAT Q64289 NEUROGENIC DIFFERENTIATION FACTOR 1; contains element LTR1 repetitive element;	ts02e12.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2227438 3' similar to SW:NDF1_RAT Q84289 NEUROGENIC DIFFERENTIATION FACTOR 1; contains element LTR1 repetitive element;	Neisseria meningitidis serogroup A strain Z2491 complete genome; segment 6/7	wd48c04.x1 Soares, NFL_T_GBC_S1 Home sapiens cDNA clone IMAGE:2331386 3' similar to gb:M37721 PEPTIDYL-GLYCINE ALPHA-AMIDATING MONOOXYGENASE PRECURSOR (HUMAN);	Campylobacter jejuni NCTC11168 complete genome; segment 4/6	y/37803.s.1 Soares fetal liver spleen 1MFLS Homo sapiens cDNA clone IMAGE:129004.3' similer to gb:X12517 U1 SMALL NUCLEAR RIBONUCLEOPROTEIN C (HUMAN);	ye82a07.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:124212.5	MR0-HT0166-181199-003-d12 HT0166 Homo sapiens cDNA	D.melanogaster mRNA for alpha 1,2 mannosidase (Berlin)	D.melanogaster mRNA for alpha 1,2 mannosidase (Berlin)	602014422F1 NCI_CGAP_Brn64 Homo saplens cDNA clone IMAGE:4150396 5	HYPOTHETICAL 75,2 KD PROTEIN C11C11.02 IN CHROMOSOME II	RC5-ET0082-310500-021-F10 ET0082 Hamo sapiens cDNA	RC5-ET0082-310500-021-F10 ET0082 Homo sapiens cDNA	S. occidentalis INV gene for invertase (EC 3.2.1.26)	Lontra canadensis cytochrome b (cytb) gene, mitochondrial gene encoding mitochondrial protein, complete	cds	GREEN-SENSITIVE OPSIN (GREEN CONE PHOTORECEPTOR PIGMENT) (KFH-G)	GREEN-SENSITIVE OPSIN (GREEN CONE PHOTORECEPTOR PIGMENT) (KFH-G)	VON WILLEBRAND FACTOR PRECURSOR (VWF)	Homo sapiens PHEX gene	wr58b09.x1 NCI_CGAP_Ut1 Homo sapiens cDNA clone IMAGE:2491865 3'
Top Hit Database Source	EST_HUMAN	Ę	Ę	EST_HUMAN	EST HUMAN	LN	EST_HUMAN	N T	EST_HUMAN	EST_HUMAN	EST_HUMAN	NT	ΝΤ	EST_HUMAN	SWISSPROT	EST_HUMAN	EST_HUMAN	N		Į.	SWISSPROT	SWISSPROT	SWISSPROT	٦	EST_HUMAN
Top Hit Acesslon No.	1 AIB62398.1	1 AF207550.1	2.6E-01 AE001811.1	2.6E-01 AIS82557.1	AI582557.1	2.6E-01 AL162757.2	2.6E-01 AI914380.1	2.6E-01 AL139077.2	2.6E-01 R10365.1	R02411.1	BE144331.1	X82641.1	X82641.1	BF343588.1	010199	BE830339.1	BE830339.1	X17604.1		AF057121.1	P87366	P87366	Q28295	Y10196.1	2.6E-01 AI978681.1
Most Similar (Top) Hit BLAST E Value	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.8E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01		2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01	2.6E-01
Expression Signal	0.81	99.0	2.36	1.89	1.89	1.05	0.97	96'0	1.6	1.14	1.18	0.67	0.67	2.99	1.89	4.49	4.49	96'0		0.5	0.93	0.93	0.5	0.91	0.51
ORF SEQ ID NO:		31289		31732	31733	31948	32512		33118	33240					33810	34080		34762			35265				
SEQ ID NO:	18386	18568	25113	18953	18953	19152	19673	24783	20229	20334			20625		20890	21168	21166	21811						22897	22994
Probe SEQ ID NO:	5760	5947	6221	6348	6348	6554	7103	7457	7721	7791	7845	8083	8083	8278	8349	8627	8627	8886		9654	9782	9782	10093	10403	10500

Page 80 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Mus musculus neuronal apoptosis inhibitory protein 6 (Naip6) gene, complete cds; and Naip3 gene, exons 2-9 Homo sapiens ATP synthase, H+ transporting, mitochondrial F1 complex, delta subunit (ATP5D), nuclear Homo sapiens ATP synthase, H+ transporting, mitochondrial F1 complex, delta subunit (ATP5D), nuclear Homo sapiens hyperpolarization activated cyclic nucleotide-gated potassium channel 4 (HCN4) mRNA Mus musculus ICR/Swiss glyceraldehyde 3-phosphate dehydrogenase (Gapd-S) gene, complete cds Ureaplasma urealydicum section 57 of 59 of the complete genome Homo sapiens Ne/K-ATPase gamma subunit (FXYD2) gene, complete cds, alternatively spliced ye11g07.r1 Stratagene lung (#897210) Homo sapiens cDNA clone IMAGE:117468 5' Botrytis cineres strain T4 cDNA library under conditions of nitrogen deprivation Mus musculus protein-L-isoaspartate (D-aspartate) O-methyltransferase 1 (Pcmt1) Homo sapiens inosital polyphosphate 1-phosphatase (INPP1) gene, complete cds 801511052F1 NIH_MGC_71 Homo sapiens cDNA clone IMAGE:3912612 5 zs11a12.r1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:684862 ! Human lambda-immunoglobulin constant region complex (germline) Arabidopsis thaliana DNA chromosome 4, contig fregment No. 29 Top Hit Descriptor EST385464 MAGE resequences, MAGM Homo sapiens cDNA Thermotoga manitima section 25 of 136 of the complete genome PM4-CT0400-310700-005-408 CT0400 Homo sapiens cDNA PM4-CT0400-310700-005-d08 CT0400 Homo sapiens cDNA Cavia cobaya mRNA for serine/threoine kinase, complete cds gene encoding mitochondrial protein, mRNA Starfish (P.ochraceus) cytoplasmic actin gene, complete cds A-AGGLUTININ ATTACHMENT SUBUNIT PRECURSOR Aquifex aeolicus section 7 of 109 of the complete genome MOLT-INHIBITING HORMONE PRECURSOR (MIH) Danio rerlo peptide YY precursor gene, complete cds CELL DIVISION PROTEIN FTSW HOMOLOG gene encoding mitochondrial protein, mRNA HYPOTHETICAL PROTEIN MG039 Mus musculus Jerky (Jrk), mRNA and 11-16 HUMAN HUMAN HUMAN HUMAN HUMAN EST_HUMAN Top Hit Database SWISSPROT SWISSPROT SWISSPROT SWISSPROT Source EST EST 몺 EST EST. F 눋 ż Ë Ħ 뉟 4502296 NT 4502296 NT 4885406|NT 10190655 6679216 Top Hit Acession 2.5E-01 AW973471.1 2.6E-01 BE883491.1 2.6E-01 AE001713.1 2.6E-01 AF141325.2 BE696604.1 2.5E-01 AE000675.1 AA251987.1 2.5E-01 AF233875.1 2.5E-01 AL161517.2 2.5E-01 AF242431.1 AF316896. AL115624. ģ 2.5E-01 U09964.1 2.6E-01 D88425.1 T89837.1 X51755. P32323 2.6E-01 P48280 2.6E-01 P47285 2.6E-01 2.5E-01 2.5E-01 2.5E-01 2.5E-01 2.5E-01 2.6E-01 2.5E-01 2.5€-01 2.5E-01 .5E-01 2.55-01 2.5E-01 2.5E-01 (Top) Hit BLAST E Most Simila 11.59 12.93 4.06 1.34 1.36 8 1.77 1.02 1.03 6.09 1.49 0.59 1.47 0.87 Expression Signal 29747 28675 36809 30996 25407 25407 25994 26274 26689 27067 ORF SEQ 27873 Ö N O 13763 15019 17303 17438 13480 14158 15454 SEQ ID 23752 24165 24991 24556 24612 24641 12921 12921 15454 15100 15104 16066 16191 16207 16735 16994 24666 ÿ 4860 1098 1766 2452 2536 2540 SEQ ID 11300 12042 12396 3459 3587 3603 4143 12478 12567 262 263 276 865 1566 1927 1927 11973 4409

Page 81 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

ORF SEQ Expression (Top) Hit Acession Database ID NO. Signal BLAST E Source Source	29896 4.69	2832 2.82	Mus musculus annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine and an annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine and an annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine and an annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine and an annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine and an an annexin V gene, intron 4 segment containing 5'LTR and gag portion of MuERV-L (murine an an an an an an an an an an an an an	29976 0.61 2.5E-01 BE896785.1 EST HUMAN	0.65 2.5E-01 M28501.1 NT	30576 12.86 2.5E-01	0.84 2.5E-01 AL163207.2 NT	22 32138 0.83 2.5E-01 AJ251973.1 NT Homo sapiens partial steerin-1 gene	82266	1.35 2.5E-01 AF134119.1 NT	33025 4.48 2.5E-01 AL183282.2 NT	33236 2.31 2.5E-01 BF109040.1 EST_HUMAN	33248 0.7 2.5E-01 BE980712.1 EST_HUMAN	33625 2.2 2.5E-01 BF038595.1 EST_HUMAN	33798	34029 3.03 2.5E-01 H53236.1 EST_HUMAN	34271 0.88 2.5E-01 M88626.1 NT	11 34909 15.98 2.5E-01 U89651.2 NT Homo sapiens match metalloproteinase MMP Rasi-1 gene, promoter region	34910 15.98	34897 2.09 2.5E-01 AF085164.1	34898 2.09 2.5E-01 AF085164.1 NT	99'1 96756	xg40c10.x1 NCI_CGAP_Uff Homo sepiens cDNA clone IMAGE:2830034.3' similar to contains Alu repetitive	35941 131 25E-01 X58401 1 NT	35962 2.03 2.5E-01 AL161505.2 NT	35963 2.03 2.5E-01 AL161505.2 INT	36495 4.3 2.5E-01 D50914.1 NT	37153 5.29	10.13 2.5E-01 AL161541.2 NT Arabidopsis thaliana DNA	25687 1.67 2.4E-01 AA936316.1 EST HUMAN	26014 2.4E-01 BF576124.1 EST_HUMAN 602132442F1 NIH_MGC
!	29896	29927				30578			82266		33025		33248	33625	33798	34029	34271	34809	34910		34898	35496	35937	35941	35962	35963	36495	37153			
e Exam ID SEQ ID NO:	69 17445	17471	4924 17499			29 18161		38 19332	7380 10014		L		97 20340	68 20709	36 20877			9435 21961	9435 21961		92 21948	10 22505	22830	L	L		55 23470	12 24122	40 25075		881 13495
Probe SEQ ID NO:	4869	4896	9	4959	5262	5529	6114	6738	۲	7413	7832	7786	7197	8168	8336	8571	8	9	94	9492	9492	10010	10438	10439	10459	10459	10955	11712	11740	2	8

Page 82 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

					5		
Probe SEQ ID NO:	Exan SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
1347		26464	21.36	2.4E-01	AJ289880.1	M	Homo sapiens KIAA0851 gene (partial), XT3 gene and LZTFL1 gene
1347	13942			2.4E-01	AJ289880.1	NT	Homo sapiens KIAA0851 gene (partial), XT3 gene and LZTFL1 gene
1427	14020	26548	0.93	2.4E-01	Y17293.1	Z	Homo sapiens FLI-1 gene, partial
1891			27.27	2.4E-01	AF267753.1	Z.	Mesembryanthemum crystallinum putative potassium channel protein Mkt1p mRNA, complete cds
1944	14528	27084	1.17	2.4E-01	AF251708.1	ΤN	Zaccys dhumnades fructose-1,6-bisphosphatase mRNA, complete cds
0700	14850	02626	07.7	F0 = F C	+ 030CF214	MAAA HOO	wg78d05.x1 Soares_NSF_F8_9W_OT_PA_P_S1 Homo sapiens cDNA clone IMAGE:2371017.3' similar to
2183			2	2.4E-01	AF111188 2	NAMOR 101	In Course Course Intervention Franciscon and Income complete additional control of the Course control of the C
2213			2	2.4E-01	P45384	TOGGSSIMIS	MANINOCIONALIN A DEDOTEASE DECLIDADO (10 4 DEDOTEASE)
2302	ı	27451	1.78	2.4E-01	801	NT	Aduitex section 12 of 109 of the complete genome
	ł	1					7h23d04.x1 NCI_CGAP_Co16 Homo sapiens cDNA clone IMAGE:3316807 3' similar to SW:PRSB_XENLA
2425				2.4E-01	BF002171.1	EST_HUMAN	042586 26S PROTEASE REGULATORY SUBUNIT 6A;
2575	15138		3.05	2.4E-01	236534.1	TN	D.discoideum (Ax3-K) panA gene
2790		27913		2.4E-01	X71783.1	NT	S.pombe swild gene
2812	15364		3.88	2.4E-01	AF030154.1	N	Bovine adenovirus 3 complete genome
3166	15780		3.27	2.45-01	U72726.1	IN	Orza longistaninata recentra kinasa-like nrotein family member D and ratroff (nan/ww) name nominista nde
3182	Ĺ	28287	1.38		X74209.1	Z	H.sapiens AGT gene, Pstl fragment of intron 4
3724	16325	28792	1.28		AF169793.1	Σ	Podospora anserina HET-C protein (Het-c) gene, complete cds
3824	į	28886	0.83		AE000312.1	NT	Escherichla coli K-12 MG1655 section 202 of 400 of the complete genome
4103			9.0	2.4E-01	D29960.1	Į.	Rattus norvegicus mRNA for alphaB crystallir-related protein, complete cds
5008					AE000305.1	M	Escherichia cdi K-12 MG1655 section 195 of 400 of the complete genome
5220		30203		2.4E-01	BE737592.1	EST_HUMAN	801572862F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:3839775 5/
5302			1.55	2.4E-01	K02402.1	TN	Human coagulation factor IX gene, complete cds
5853	- 1			2.4E-01	A1925707.1	EST_HUMAN	wo33d05.x1 NCI_CGAP_Gas4 Homo sapiens cDNA clone IMAGE:2457129 3'
5653	- 1		0.83	2.4E-01	AI925707.1	EST_HUMAN	wo33d05.x1 NCL_CGAP_Gas4 Homo sapiens cDNA clone IMAGE:2457129 3'
5676				2.4E-01	D50871.1	NT	Glycine max mRNA for mitotic cyclin b1-type, complete cds
5836	18460	31182	7.92		AF091216.1	IN	Mus musculus Wrn protein (Wrn) gene, complete cds
5836		31183		2.4E-01	AF091218.1	NT	Mus musculus Wrn protein (Wm) gene, complete cds
6050	24754		1.02		AJ133838.2	NT	Branchlostoma floridae mRNA for calmodulin 2 (caM2 gene)
							7154004.x1 NCI_CGAP_Br16 Homo sapiens cDNA clone IMAGE: 3338503 3' similar to SW: SFR4_HUMAN
6054	18672	31411	2.36	2.4E-01	BF592336.1	EST HUMAN	
6138	18752	31510	2.5	2.4E-01	AF035546.1	NT	Drosophila melanogaster p38a MAP kinase gene, complete cds

Page 83 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Homo sapiens HSPC142 protein (HSPC142), mRNA	AV733787 cdA Homo sapiens cDNA clone cdAADE11 5'	wc62c11.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2323220 3' similar to gb:.J03484 PROCOLLAGEN ALPHA 2(1) CHAIN PRECURSOR (HUMAN);	Bos taurus guanylyl cyclaso-activating protein 2 (guca2) mRNA, complete cds	Mus musculus DXImx48e protein (DXImx48e) mRNA, complete cds	Streptococcus pneumoniae rr08 and hk08 genes; two component system 08	Streptococcus pneumoniae rr08 and hk08 genes; two component system 08	Tetrahymena thermophila macronuclear gene encoding ribosomal protein L3, excns 1-2	601877679F1 NIH_MGC_55 Homo sapiens cDNA clone IMAGE:4106298 5'	602086188F1 NIH_MGC_83 Homo sapiens cDNA clone IMAGE:4250372 57	Campylobacter jejuni NCTC11168 complete genome; segment 4/6	Campylobacter jejuni NCTC11168 complete genome; segment 4/6	wd43e02.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2330906 3' sImilar to contains	MER22.b1 TAR1 repetitive element;	Drosophila melanogaster SKPB gene, complete cds	Drosophila melanogaster SKPB gene, complete cds	COLLAGEN ALPHA 1(X) CHAIN PRECURSOR	Arabidopsis thatiana DNA chromosome 4, contig fragment No. 6	Mus musculus type 1 sigma receptor gene, complete cds	P. asiatica mosaic virus genomic RNA	Homo sapiens fragile 16D oxido reductase (FOR) gene, exan 6	Arabidopsis thaliana ethylene-insensitive3-like1 (EIL1) mRNA, complete cds	Mus musculus mRNA for putative mc7 protein (mc7 gene)	Gallus gallus gene coding for e-actin	RC3-CT0413-100800-023-b06 CT0413 Homo capiens cDNA	Homo sapiens chromosome 21 segment HS21C081	aromatasa (Poephila guttata=zabra finches, ovary, mRNA, 3188 nt)	Mycoplasma genitalium section 35 of 51 of the complete genome	Methanococcus jannaschil section 138 of 150 of the complete genome	601142073F1 NIH_MGC_14 Homo sapiens cDNA clone IMAGE:3505818 5	Brassica napus sig gene for S-locus glycoprotein, cultivar T2	Mus musculus cdh5 gene, exon 1, pertial	Homo sapiens partial intron 3 of the wild type AF-4/FEL gene
Top Hit Database Source		EST_HUMAN /	EST HUMAN		E	F	LZ.	Ę	EST HUMAN	EST_HUMAN	Г	Ę		EST_HUMAN		TN I	SWISSPROT	Į,	Z	Z		Į,		IN	EST_HUMAN	Ę	Į.	IN	LN	EST_HUMAN	E E		TN
Top Hit Acession No.	7681801 NT	AV733787.1	A1698989.1	L43001.1	AF229644.1	AJ006397.1	AJ006397.1	AJ012585.1	BF242794.1	BF678275.1	AL139077.2	AL139077.2		AI693515.1		AF220067.1	Q03692	AL161494.2	AF030199.1	221647.1	AF217491.1	AF004213.1	AJ278191.1	V01507.1	BF229975.1	AL163281.2	S75898.1	U39713.1	U67596.1	BE311893.1	AJ245480.1	Y10887.2	AJ235353.1
Most Similar (Top) Hit BLAST E Vatue	2.4E-01	2.4E-01		2.4E-01	2.4E-01		2.4E-01		2.4E-01	2.4E-01	2.4E-01	2.4E-01 /			2.4E-01	2.4E-01	2.4E-01 (2.4E-01	2.4E-01	2.4E-01		2.4E-01		2.4E-01	2.4E-01	2.4E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01
Expression Signal	2.28	8.0	2.43	8.8	1.06	17.0	17.0	1.66	26.0	0.58	0.58	0.58		6.84	9.0	9.0	1.95	3.25	2.9	2.28	1.91	2.65	2.02	2.18	1.5	2.31	16:0	4.4	17.02	3.44	1.19	2.75	1.28
ORF SEQ ID NO:	31619	31669	32055	32772	33108	33591	33592	33752	33994		34526	34527		34826	35083	35084	35823	36192	36260		37145						25538		25803	28092	28774	26800	
Exon SEQ ID NO:	1	18898	19252	l	20218			Ε	21074	21127	21598	21596	ı	- 1		22120		23179	23243			24853				24662	13047		13318	13580	14239		14669
Probe SEQ ID NO:	6240	6290	9999	7381	7709	8139	8139	8290	8535	8288	9059	6906		9482	9620	8620	10335	10647	10715	11081	11665	11807	11866	12086	12320	12562	412	965	695	696 6	1647	1674	2089

Page 84 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	601175562F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:3531015 5'	Human erythropoietin gene, complete cds	Mariniabilia agarovorans gyrB gene for DNA gyrase subunit B, partial cds, strain:IFO 14957	no16d06.s1 NCI_CGAP_Phe1 Homo sepiens cDNA clone IMAGE:1100843 3' similar to contains Alu repetitive element; contains element THR repetitive element;	yh21b07 s1 Soeres placenta Nb2HP Homo sapiens cDNA clone IMAGE:130357 3'	yr97h10.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:213283 5'	GSTA5=glutathione S-transferase Yc2 subunit {5' region, intron 1} [rats, Morris hepatoma cell line, Genomic, 2212 nt, segment 1 of 3]	Homo sapiens KIAA0450 gene product (KIAA0450), mRNA	y17f01.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:149017 5'	Mus musculus renin (Ren-1c) gene, promoter region	Synechocystis sp. PCC6803 complete genome, 1/27, 1-133859	Homo sapiens mitogen-activated protein kinase p38detta (PRKM13) mRNA, complete cds	Homo sapiens nuclear transport factor 2 (placental protein 15) (PP15) mRNA	Human phenylethanolamine N-mathyltransferase gene, complete cds	Mus musculus tulip 1 mRNA, complete cds	Escherichia coli K-12 MG1655 section 130 of 400 of the complete genome	Homo sepiens mRNA for KIAA1512 protein, pertial cds	7k30b06.x1 NCI_CGAP_Ov18 Homo sapiens cDNA clone IMAGE:3476699 3' similar to SW:CAG_SMSAV P03330 GAG POLYPROTEIN [CONTAINS: CORE PROTEIN P15; INNER COAT PROTEIN P12; CORE CHELL DEOTEIN P30: NILCI EOPROTEIN P401	Cfamiliaris rom 1 cana	Vittaforma corneum small subunit ribosomal RNA gene	23S rRNA [Leuconostoc carnosum, Genomic, 2866 nt]	as27e12.x1 Barstead aorta HPLRB6 Homo sapiens cDNA clone IMAGE:2318446 3' similar to gb:X13238 CYTOCHROME C OXIDASE POLYPEPTIDE VIC PRECURSOR (HUMAN);	es27e12.X1 Barstead acrta HPLRB6 Homo saplens cDNA clone IMAGE:2318446 3' similar to gb:X13238 CYTOCHROME C OXIDASE POLYPEPTIDE VIC PRECURSOR (HUMAN);	Oryctolagus cuniculus cytochrome oxidase subunit VIa (coxVIa2) mRNA, complete ods, nuclear gene for mitochondrial product	as42f12.x1 Barstead aorta HPLRB6 Homo sapiens cDNA done IMAGE:2319887.3' similar to contains Alu	Homo saplens hypothetical protein FLJ20345 (FLJ20345), mRNA
Top Hit Dambase Source	EST_HUMAN	I	I IN	EST HUMAN	T	EST_HUMAN			EST_HUMAN	Z	LN L	IN		N TN	LX.	LN	LN TA	NAME TOO	Т	Ł	Σ	EST HUMAN		Ę	NOT LINAN	NT
Top Hit Acession No.	BE297718.1	M11319.1	AB015033.1	AA601379.1	R21732.1	H69836.1	S82821.1	7662133 NT	R82252.1	L78789.1	D90899.1	AF092535.1	5031984 NT	J03280.1	AB032400.1	AE000240.1	AB040945.1	05050304.4	Yorks 1	39/12.1	S60371.1	AI708840.1	AI708840.1	AF109090 1	A1740440 4	8923323 NT
Most Similar (Top) Hit BLAST E Value	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01		2.3E-01	2.3E-01		2.3E-01		2.3E-01	2.3E-01		2.3E-01		2.3E-01	220	_					2.3F.01	1	_
Expression Signal	2.03	1.18	1.42	0.83	96.9	0.78	1.02	5.14	0.83	2.4	0.87	2.16	6.13	0.62	0.62	6:0	2.39	300	20.7	1.19	0.78	2.34	2.34	0.78		0.7
ORF SEQ ID NO:			26552			28507			29468		29578	29613	29690		30191	30372	30552		31050		31274	31461			\	
SEQ ID NO:	1	15236	14024	15606		16025	16507	16607	17028	17074	17131	17169	17234	17585		17961	18140	7300	⊥_	ı	1	18712	18712	10355	1	ĽĽ
Probe SEG ID NO:	2489	2678	2851	2890	3120	3417	3908	4009	4442	4489	4548	4586	4652	5180	5202	5403	2055	2624	5724	5831	5926	9609	8609	8782	908	7165

WO 01/57277

Page 85 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Secale cereale omega secalin gene, complete cds	Glycine max resistance protein LM17 precursor RNA, partial cds	Mus musculus myosin XV (Myo15), mRNA	801511573F1 NIH_MGC_71 Homo sepiens cDNA clone (MAGE:3912859 5	za12e08.r1 Soares fetal liver spleen 1NFLS Home sapiens cDNA clone IMAGE:292358 5'	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 58	Oxydriche nova macronuclear telomere-binding protein alpha subunit (tel-alpha alanine version) gene, complete cds	Mus musculus prosaposin (psap\SGP-1) gene, complete cds	EST84061 Rhabdomyosarcoma Homo sapiens cDNA 5' end similar to DneJ homolog (GB:X83368)	EST84061 Rhabdomyosarcoma Homo sapiens cDNA 5" end similar to DnaJ homolog (GB:X63368)	Mus musculus phosphatidylinositol 3-kinase catalytic subunit delta (Pik3cd), mRNA	601120110F1 NIH_MGC_20 Homo saplens cDNA clone IMAGE:2868739 5'	EST376533 MAGE resequences, MAGH Homo sapiens cDNA	Haemophilus influenzae genes for Hincli restriction-modification system (Hincli methyltransferase (EC	2.1.1.72) and Hincil endonuclease (EC $3.1.21.4$))	PM2-DT0036-281299-001-f04 DT0036 Homo sapiens cDNA	MR0-HT0559-240400-014-g11 HT0559 Homo sapiens cDNA	Rhizoblum leguminosarum partial genomic DNA for exopolysaccharide biosynthesis genes	Murine hepatitis virus strain 2, complete genome	601648155R2 NIH_MGC_59 Homo sapiens cDNA clone IMAGE:4102092 3'	Mus musculus tissue factor pathway inhibitor (TFPI) mRNA, complete cds	Mus musculus fissue factor pathway inhibitor (TFPI) mRNA, complete eds	Mus musculus partial mRNA for muscle protein 534 (mg534 gene)	Mus musculus partial mRNA for muscle protein 534 (mg534 gene)	Chlamydophila pneumoniae AR39, section 4 of 94 of the complete genome	AV709736 ADC Homo sapiens cDNA clone ADCAGH01 5'	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	HCOEST44 HT29M8 Homo sapiens cDNA clone HCoE44 5	chn1424.seq.F Human fetal heart, Lambda ZAP Express Homo sapiens cDNA 5'	PM4-SN0012-030400-001-b06 SN0012 Homo sapiens cDNA	x/21407.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2813773 3' similar to TR:Q92175	92175 LYSYL OXIDASE-RELATED PROTEIN 2. ;contains PTR5.b2 TAR1 repetitive element;
Top Hit Database Source	Ę			EST_HUMAN	Г	Γ	Ł		EST_HUMAN	EST HUMAN		EST_HUMAN	EST_HUMAN	Г	╗		EST_HUMAN		NT NT	EST_HUMAN			INT.		⇈	EST_HUMAN /	Т	EST_HUMAN	EST_HUMAN	П		EST HUMAN C
Top Hit Acession No.	01 AF000227.1	01 AF175389.1	6754779 NT	01 BE888071.1	01 N80983.1	01 AL161558.2	01 M88931.1	J57999.1	01 AA372164.1	4A372164.1	2.3E-01 6679318 NT	3E277860.1	01 AW984460.1)1 X52124.1	01 AW364633.1	2.3E-01 BE173060.1	2.3E-01 AJ293261.1	01 AF201929.1	1 BF133577.1	01 AF004833.1	01 AF004833.1	01 AJ250189.1	01 AJ250189.1	01 AE002167.2	1 AV709736.1		1 T27231.1		2.3E-01 AW863940.1		2.3E-01 AW303623.1
Most Similar (Top) Hit BLAST E Value	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01 U57999.1	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01		2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01 U45426.1	2.3E-01	2.3E-01	2.3E-01		2.3E-01
Expression Signal	0.69	2.42	3.63	1.83	2.68	0.58	1.93	9.0	0.87	28.0	9.65	0.51	0.76	,	1.22	0.55	2.6	1.93	0.94	5.86	1.85	1.85	1.85	1.85	2.49	1.6	2.82	57.94	1.31	1.61		3.1
ORF SEQ ID NO:	32721	32837		32998		33243	33387	33882	34534		34958	35112	35169		35225	35264	35326	35373			36432	36433	36619	36620	36817				-			30810
Exen SEQ ID NO:	19858	19969	20116	20121	20240	20336	20477	20970	21604	21604	22001	22144	22198		22244	22279				22845	23414	23414	23580	23580	23761	24068	24172	24228	24804	24246		25002
Probe SEQ ID NO:	7331	7445	7603	7608	7732	7793	7935	8430	2906	2906	9501	9644	9697	9, 10	9746	9781	9847	9903	10339	10351	10893	10883	11068	11068	11230	11624	11788	11876	11899	11908		11969

Page 86 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor Source	HUMAN 601507202F1 NIH_MGC_71 Homo sapiens cDNA done IMAGE 3908689 5'	HUMAN 60214449F1 NIH_MGC_48 Homo sapiens cDNA clone IMAGE:4297719 5'	Rattus norvegicus mRNA for acid gated ion channel	Pleurodakes walt distal-less like protein PwOlx-3 (PwOlx-3) mRNA, complete cds	Rattus norvegicus mRNA for acid gated ion channel	nsc39h12.x1 Lupski_sciatic_nerve Homo sapiens cDNA clone IMAGE:3395950 3' similar to contains element UMAN MER38 repetitive element;	Г		Homo sapiens PPAR delta gene, promoter region	Trimeresurus malabaricus cytb gene, partial cds; mitochondrial gene for mitochondrial product	Fresh-water sponge Emf1 alpha collagen (COLF1) gene	UMAN 602085608F1 NIH_MGC_83 Homo sepiens cDNA clone IMAGE:4249689 5'	Г	Г	Homo sapiens chromosome 21 segment HS21C018		UMAN PM2-HT0353-261289-003-a12 HT0353 Homo sapiens cDNA	Homo sapiens FRA3B common fragile region, diadenosine triphosphate hydrolase (FHIT) gene, exon 5	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 62	Xiphophorus maculatus truncated Rex1 retrotransposon reverse transcriptase (RT) pseudogene	Drosophila melanogaster UNC-119 (unc-119) gene, complete cds	Mus musculus mixed lineage kinase 3 (Mik3) and two pore domain K+ channel subunit (Kcnk6) genes,	Mis misculas MAP kinasa kinasa i Makki) mBNA Complete ode	Mus musculus MAP kinase kinase (Mekk1) mRNA complete cds	Human scRNA (BC200 beta) pseudogene	Human scRNA (BC200 beta) pseudogene	Human beta-cytoplasmic actin (ACTBP9) pseudogene	UMAN zq87c05.r1 Stratagene hNT neuron (#937233) Homo sapiens cDNA clone IMAGE:648968 5'	Mus musculus vinculin gene, excn 3	Borrelia burgdorferi (section 23 of 70) of the complete genome
Top Datat	EST_HU	EST_HU	뉟	Έ	Z	EST HUMAN	EST HUMAN	EST_HUMAN	Z	뒫	Z	EST_HUMAN	EST_HUMAN	EST_HUMAN	Ę	EST_HUMAN	EST_HUMAN	¥	Ψ	¥	LΝ	5	 	5	z	Ę	뉟	EST_HUMAN	F	Ę
Top Hit Acession No.		BF663319.1	AJ006519.1	U49645.1	AJ006519.1	BF475611.1	AA094108.1		AF187850.1	AF171901.1	M34640.1	BF677538.1	BE618258.1	BE618258.1	AL163218.2	BE155625.1	BE155625.1	AF020503.1	AL161562.2		AF119102.1	AF155142 1	Ī			U01307.1	D50604.1	AA211216.1		AE001137.1
Most Similar (Top) Hit BLAST E Value	•		2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.3E-01	2.2E-01								2.2E-01		2.2E-01			2.2E-01	2.2E-01	_					2.2€-01	2.2E-01	
Expression Signal	10.96	1.94	3.11	1.38	1.67	2.57	1.26	0.91	2.85	3.89	3.16	5.61	1.27	1.27	1.17	4.28	4.28	1.64	2.67	1.18	1.26	2 03	2.59	2.59	1.36	1.36	1.35	2.86	1.33	1.79
ORF SEQ ID NO:	30511						30710	25252	26738		27287	27586	27751	27752		27993	27994	•				20330		29379	29475	29476		7862		
Exon SEQ ID NO:	25053	24340	24369	24429	24369	24614	24888	12769	14204	14643	14714	15014	15185	15185	15260	15523	15523	15563	16047	16484	16877	16886	16937	16937	17033	17033	17527	17532	17761	17768
Probe SEQ ID NO:	12007	12057	12107	12205	12211	12480	12668	8	1611	2063	2138	2447	2623	2623	2703	2906	2908	2947	3439	3886	4291		4350	4350	4447	4447	4952	4957	5196	5203

Page 87 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

		Γ								_	<u> </u>					T	Т	٦								•	Г	Γ			
	Top Hit Descriptor	MR0-HT0067-201099-002-c10 HT0067 Homo sapiens cDNA	histamine H2-receptor (rats, Genomic, 1928 nt)	Homo saplens diaphanous (Drosophila, homolog) 2 (DIAPH2), transcript variant 156, mRNA	Synechocystis sp. PCC6803 complete genome, 19/27, 2392729-2538999	Gallus gallus T-box containing protein (Ch-TbxT) mRNA, complete cds	Gallus gallus T-box containing protein (Ch-TbxT) mRNA, complete cds	Homo sapiens gene for fukutin, complete cds	AV756238 BM Homo saplens cDNA clone BMFAHC06 5'	Streptococcus pyogenes phosphotidy/glycerophosphate synthese (pgsA) and ABC transporter ATP-binding protein (stpA) genes, complete cds; and unknown genes	Streptococcus pyogenes phosphotidy/glycerophosphata synthase (pgsA) and ABC transporter ATP-binding	protein (stpA) genes, complete cds; and unknown genes	Human glycophorin B gene, exon 4	Human glycophorin B gene, exon 4	Homo sapiens homeobox B7 (HOXB7) gene, pertial cds; and homeobox B6 (HOXB6), homeobox B6 (HOXB4) and homeobox B3 (HOXB3) perse, complete rds	March 1 to the control of the contro	Mus musculus nm23-M1 gene, promorer region	E.coll sep8 and sep8 genes	Mouse HD protein mRNA, complete cds	Mouse HD protein mRNA, complete cds	Thermotoga maritima section 25 of 136 of the complete genome	Mus mucculus ICR/Swiss glyceraldehyde 3-phosphate dehydrogenase (Gapd-S) gene, complete cds	PM3-CT0263-241289-009-b07 CT0263 Homo sapiens cDNA	Mus musculus deformed epidermal autoregulatory factor 1 (Drosophila) (Deaf1), mRNA	MR1-TN0045-110900-006-c02 TN0045 Hamo sapiens cDNA	za04f08.r1 Soares melanocyte 2NbHM Homo sapiens cDNA clone IMAGE:291591 5'	LARGE PROLINE-RICH PROTEIN BAT2 (HLA-B-ASSOCIATED TRANSCRIPT 2)	Xenopus laevis mRNA for kinesin-like protein 3 (xklp3)	Mus musculus osteoblast specific factor 2 (OSF-2), mRNA	Brachydanio rerio ependymin beta and gamma chains (Epd) gene, complete cds	CYCLIC NUCLEOTIDE GATED CHANNEL, ROD PHOTORECEPTOR, ALPHA SUBUNIT (CNG CHANNEL 3) (CNG-3) (CNG3)
Top Hit	Database Source	EST_HUMAN	Z	FN	LZ.	FZ	LΝ	LN	EST_HUMAN	, t		NT	FZ	Į.	Ļ		Z	NT	NT	NT	NT	Þ	EST_HUMAN	Ę	EST_HUMAN	EST_HUMAN	SWISSPROT	¥	N FA	IN	SWISSPROT
	No.	2.2E-01 BE141035.1		5803002 NT	2.2E-01 D64000.1		1 U67087.1		2.2E-01 AV756238.1	2.2E-01 AF082738.1		1		2.2E-01 M24136.1	14 AE287087 4						2.2E-01 AE001713.1	2.2E-01 U09964.1	2.2E-01 AW855039.1	8393247 NT	2.2E-01 BF376354.1	W02988.1	P48634		657428	01 M89643.1	090380
Most Similar	(10p) nit BLASTE Value	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2€-01	2.2€-01	2.2€-01	2.2E-01		2.2E-01	2.2E-01	2.2E-01	2 2E-04	2.25-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2€-01	2.2E-01	2.2E-01	2.2E-01	2.2E-01	2.2€-01	2.2E-01 Q90980
	Signal	1.2	6.0	2.46	3.53	0.73	0.73	0.85	9.14	1.46		1.46	2.01	2.01	89.0	80.0	3.06	0.84	0.57	0.57	3.48	1.02	3.12	1.82	1.95	1.24	14.03	0.74	17.0	3.69	0.59
	ON OO OO OO	30277		31267		31525	31526	32212	32503	32562			32723	32724	380EE	COACC		33479	34277	34278	34290	34310		34509	34604	34673	34903	34750			
Exon	SEQ ID NO:	17853	l	18541	18552	18763	18763	19398		19715	l	1	19860		20107	1	20202	- 1			21366	21386		21580		21730	L	L		21898	L
Probe	SEQ ID NO:	5291	5316	5919	2930	6150	6150	6807	7093	7183		7183	7333	7333	7688	3 2	3	8032	8815	8815	8827	8847	8952	9043	9128	9213	9231	9274	9285	9288	9539

Page 88 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Page 89 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	IMMEDIATE-EARLY PROTEIN IE180	Orchestia cavimana calcium-binding protein BP23 precursor (BP23) gane, complete cds	Homo sapiens mRNA for KIAA1215 protein, partial cds	Ното sapiens pshsp47 gene, complete cds	Lycopersicon esculentum homeobox 1 protein (THox1) mRNA, partial cds	Vampire bat (D.rotundus) plasminogen activator mRNA, complete cds	602152001F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4293001 5	Doto fragilis mitochondrial 16S rRNA gene, partial	Human olfaciory receptor (OR17-2) gene, partial cds	VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KV3.3 (KSHIIID)	VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KV3.3 (KSHIIID)	Archeeoglobus fulgidus section 135 of 172 of the complete genome	Canis familians keratin (KRT9) gene, complete cds	Gycine max matate dehydrozenase (Mdh-2) gene, nuclear gene encoding mitochondrial protein, partial cda	Chains may mainte debt of consequent (Math. 2)	vigation make unique general (works) gene, increasing encounting minoring protein, perusing on vigation of the contraction of t	TOTAL OF THE STREET THE PROPERTY OF THE PROPERTY COLOR OF THE PROPERTY OF THE	Mus musculus erythrocyte protein band 4.1-like 3 (Epb4.1B), mRNA	Heemophilus influenzae hmcD, putative heemocin processing protein (hmcD), putative ABC transporter (hmcB), putative heemocin structural protein (hmcA), and haemocin immunity protein (hmcI) genes, complete	DKFZp434H0614 r1 434 (synonym: rkes3) Hano sapiens cDNA clane DKFZp434H0614 5'	DKFZp434H0614_r1 434 (synonym: ktes3) Hamo sapiens cDNA clone DKFZp434H0614 5'	Homo sapiens APCL gene, exon 9	S.cerevisiae chromosome II reading frame ORF YBL025w	1y/11e10.r1 Soares melanocyte 2NbHM Homo sapiens cDNA clone IMAGE:270954 5	3y11e10.r1 Sogres melanocyte 2NbHM Homo sapiens cDNA clone IMAGE: 270954 5	A.thaliana mRNA for AtRanBP1b protein	Homo sapiens p53R2 gene for ribonucleotide reductase, exon 6	Beta wilgaris mRNA for elongation factor 1-beta	DIACYLGLYCEROL KINASE, DELTA (DIGLYCERIDE KINASE) (DGK-DELTA) (DAG KINASE DELTA) (80 KD DIACYLGLYCEROL KINASE)	602131427F1 NIH_MGC_81 Homo sapiens cDNA clone IMAGE:4270831 5
Top Hit Deterberse Source	SWISSPROT	¥	N T	N.	LN	Z	EST_HUMAN	Į.	Į.	SWISSPROT	SWISSPROT	N.	ΝŢ	۲	ŀ	EST HIMAN	-1	LN.		EST HUMAN	EST HUMAN	N	Ę	EST_HUMAN	EST_HUMAN	NT	ΙN	LN	SWISSPROT	EST_HUMAN
Top Hit Acesslon No.	P11675	2.1E-01 AF124526.1	2.1E-01 AB033041.1	2.1E-01 AB010273.1	2.1E-01 U78409.1	2.1E-01 J05082.1	2.1E-01 BF672695.1	2.1E-01 AJ223392.1	2.1E-01 U04642.1	Q01956	001956	AE000972.1	2.1E-01 AF000949.1	1 AF068687.1	1 A Eneses 7 4	TR7354 1		7305030 NT	1 169300 4	AL040537.1	1 AL040537.1	AB022524.1	Z35786.1	N42536.1	N42536.1	X97378.1	AB036529.1	297067.1	P52824	BF574254.1
Most Similar (Top) Hit BLAST E Value	2.1E-01 P11875	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01 Q01956	2.1E-01 Q01956	2.1E-01	2.1E-01	2.1E-01	2.15.04	2 1F-01		2.1E-01	, C	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01	2.1E-01
Expression Signal	1.22	1.38	1.51	1.83	1.63	0.98	8.55	1.18	2.04	1.24	1.24	2.17	2.02	1.14	4.	89 0		1.19	4 02	0.82	0.82	0.47	5.93	9.0	9.0	2.95	1.57	1.04	1.96	29'0
ORF SEQ ID NO:	29177					96506		32368	86ZZE	32825	32826		33088	33130	33434				33804	34190	34191		34426	34880	34881			35719	35745	35751
Exan SEQ ID NO:					17656					l			20201	20239		20273	П	П	20979	l								22727	22758	22764
Probe SEQ ID NO:	4129	4336	4465	4878	5083	5434	5504	6967	6379	7436	7436	7447	7692	7731	7731	7785	6	È	8439	8732	8732	8888	8967	9423	9423	9432	9538	10232	10263	10269

Page 90 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

				;			
Probe SEQ ID NO:	Exon SEQ ID NO:	ORF SEQ ID NO:	Expression Signal	Most Similar (Top) Hit BLAST E Value	.Top Hit Acession No.	Top Hit Database Source	Top Hit Descriptor
10505	22999	36007	0.5	2.1E-01	AF294296.1	ħ	Anolis lineatopus isolate NG NADH dehydrogenase subunit 2 (ND2) gene, complete cds; mitochondrial gene for mitochondrial product
11438	23888		2.24	2.1E-01	11036847 NT	NT	Homo sapiens pancreatic polypeptide 2 (PPY2), mRNA
11451	23901	69698	2.34	2.1E-01	BE180422.1	EST_HUMAN	RC3-HT0622-040500-013-b11 HT0622 Homo sapiens cDNA
11841	24602		1.39	2.1E-01	X57624.1	Ŋ	Drosophila melanogastar ALA-E8 DNA, repeat region
12183	24418		1.48	2.1E-01	AF217490.1	Z	Homo sapiens fragile 16D oxdo reductase (FOR) gene, exons 8, 9, and partial cds
12485	24593		1.72	2.1E-01	BE622149.1	EST_HUMAN	601440712F1 NIH_MGC_72 Homo sepiens cDNA clone IMAGE:3915675 5'
12607	24691	30858	2.08	2.1E-01	BE672330.1	EST_HUMAN	7a59e02.x1 NOL CGAP_GC8 Homo sepiens cDNA clone IMAGE:3223034 3'
12812	24695	30861	1.28	2.1E-01	AJ276505.1	LN IN	Mus musculus genomic fragment, 279 Kb, chromosome 7
214	12875	25362	1.86	2.0E-01	AB017437.1	NT	Gallus gallus mRNA for avena, complete cds
529	13190		2.2	2.0E-01	1095027	IN	Homo sapiens CGI-18 protein (LOC51008), mRNA
728	13348		1.24	2.0E-01	M77085.1	TN	O.cunniculus germline IgH heavy chain V-H pseudogene, allotype VHa2
88 83	ı	25968	1.76		AF027865.1	LN	Mus musculus Major Histocompatibility Locus class II region
1049	13656	26167		2.0E-01	D90905.1	IN	Synechocystis sp. PCC6803 complete genome, 7/27, 781449-920915
1164	13768	26276	3.24	2.0E-01	AL163213.2	ΙN	Homo sapiens chromosome 21 segment HS21 C013
1297	13891	26414		2.0E-01	AJ132695.5	Z	Homo sapiens rac1 gene
1351	13946	26470	ZZ'1	2.0E-01	AW384937.1	EST_HUMAN	PM1-HT0422-291289-002-c06 HT0422 Homo sapiens cDNA
1507	14099		1.22	2.0E-01	AJ243957.1	LN	Pium pox virus strain M, complete genome, isolate PS
1534	14126	26663	23.08	2.0E-01	4503408 NT	IN	Homo sapiens dystrobrevin, alpha (DTNA), mRNA
1599	14191	26722	3.03	2.0E-01	AB007974.1	Ę	Homo sapiens mRNA, chromosome 1 specific transcript KIAA0505
1604	14196		1.23	2.0E-01	AF260700.1	Į.	Homo sapiens sodium/todide symporter mRNA, partial cds
1735		26868	1.17	2.0E-01	U22346.1	ĮN.	Human bradykinin B1 receptor (bradyb1) gene, complete cds
1755	14345		1.83	2.0E-01	AF111170.3	IN	Homo sapiens 14q32 Jagged2 gene, complete cds; and unknown gene
1795	14385		1.99	2.0E-01	U67525.1	IN	Methanococcus jannaschii section 67 of 150 of the complete genome
1934	14518		1.14	2.0E-01	BE871330.1	EST_HUMAN	601449441F1 NIH_MGC_65 Hamo sapiens cDNA clane IMAGE:3853330 5
1934	14518	27074	1.14	2.0E-01	BE871330.1	EST_HUMAN	601449441F1 NIH_MGC_65 Homo sapiens cDNA clone IMAGE:3853330 5'
1937	14521	27077	1	2.0E-01	TN 8522238	IN	Homo sapiens hypothetical protein FLJ10120 (FLJ10120), mRNA
2386	14955		1.64	2.0E-01	X82877.1	LN	H.sapiens Na+-D-glucose cotransport regulator gene
2915	15532		99'0	2.0E-01	AF074990.1	LN	Homo sapiens full length insert cDNA YH85A11
3534	16139	28621	0.7	2.0E-01	P46807	SWISSPROT	HOMEOBOX PROTEIN GLÁBRAZ (HOMEOBOX-LEUCINE ZIPPER PROTEIN ÁTHB-10) (HD-ZIP PROTEIN ATHB-10)
		\					xp15b02.x1 NCI_CGAP_HN9 Homo sapiens cDNA clone IMAGE:2740395 3' similar to contains element
3626	-1	`.		2.05-01	AW Z38005.1	ESI_HUMAN	MEK21 repositive element;
3768	16369	28835	0.8	2.0E-01	P34641	SWISSPROT	CED-11 PROTEIN

Page 91 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Sus scrofa	C.parasitica eapC gene	Mus musculus neuronal apoptosis inhibitory protein 6 (Naip6) gene, complete cds; and Naip3 gene, exons 2-9 and 11-16	QV4-EN0032-190500-223-e03 EN0032 Homo sapiens cDNA	Homo sapiens hypothetical protein ASH1 (ASH1), mRNA	HOMEOBOX PROTEIN GLABRA2 (HÖMEÖBOX-LEUCINE ZIPPER PROTEIN ATHB-10) (HD-ZIP PROTEIN ATHB-10)	Rat SOD-2 gene for manganese-containing superoxide dismutase	Homo sepiens dual oxidase-tike domains 2 (DUOX2), mRNA	Frubripes DNA encoding for valyI-tRNA synthetase	Saccharomyces cerevisiae Hal5p (HAL5) mRNA, complete cds	Human hepatocyta growth factor gene, exon 1	Mauratus mu class glutathione transferase gene	PM1-CT0247-141099-001-g06 CT0247 Homo sapiens cDNA	Mycoplasma genitalium section 46 of 51 of the complete genome	Mus musculus phosphofructokinase-1 C isozyme (Pfkc) gene, exans 3 through 7	Homo sapiens mRNA for FLJ00016 protein, partial cds	Andes virus strain Ol23133 glycoprotein G1 and G2 precursor, gene, partial cds	M.musculus scp2 gene exon 14	601344648F1 NIH_MGC_8 Homo sapiens cDNA clone IMAGE:3877794 5	Dictyostelium discoldeum random slug cDNA19 protein (rsc19) mRNA, partial cds	Arabidopsis pyruvate decarboxylase-2 (Pdc2) gene, complete cds	Chlamydia trachomatis section 5 of 87 of the complete genome	DAUGHTERLESS PROTEIN	DAUGHTERLESS PROTEIN	Homo saplens filamin 2 (FLN2) mRNA, complete cds	Arabidopsis thaliana root gravitropism control protein (PIN2) gene, complete cds	Arabidopsis thaliana root gravitopism control protein (PIN2) gene, complete cds	Homo sapiens cAMP specific phosphodiesterase (PDE4C) gene, exons 2 through 12	Homo sapiens cAMP specific phosphodiesterase (PDE4C) gene, exons 2 through 12	D.melanogaster DNA mobile element (hoppet)	R.norvegicus mRNA for NTR2 receptor	Salvelinus pluvius mRNA for transfertin, complete cds
Top Hit Database Source	Z,	۲	LΖ	EST HUMAN	F	SWISSPROT	NT	Į	ΝT	N _T	۲	Σ	EST_HUMAN	IN	LΝ	ΙN	LΝ	ĮΝ	EST_HUMAN	IN	'n	ΙN	SWISSPROT	SWISSPROT	NT.	Į,	IN	LN	TN	TN	NT	ŊŢ
Top Hit Acession No.	2.0E-01 Z46906.1	2.0E-01 X83997.1	2.0E-01 AF242431.1	2.0E-01 BE826165.1	8922080 NT	246607	2.0E-01 X56600.1	11432540 NT	2.0E-01 X91856.1	2.0E-01 U15300.1	2.0E-01 M75967.1	X61033.1	4W360865.1	J39724.1	2.0E-01 AF250371.1	2.0E-01 AK024427.1	2.0E-01 AF028028.1	2.0E-01 X91151.1	2.0E-01 BE562247.1	J82511.1	2.0E-01 U71122.1	2.0E-01 AE001278.1	211420	1 P11420	2.0E-01 AF146692.1	2.0E-01 AF086907.1	2.0E-01 AF086907.1	2.0E-01 AF157814.1	2.0E-01 AF157814.1	2.0E-01 X78388.1	X97121.1	2.0E-01 D89088.1
Most Similar (Top) Hit BLAST E Vatue	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01 P46607	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01 X61033.1	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01 P11420					2.0E-01	2.0E-01	2.0E-01	2.0E-01	2.0E-01
Expression Signal	0.78	0.68	0.76	8.43	7.09	0.62	2.38	2	0.69	6.48	0.71	3.84	3.63	0.68	1.18	1.53	6.45	2.91	0.53	1.03	0.85	4.35	0.51	0.51	1.98	1.79	1.79	0.53	0.53	0.72	0.88	2.77
ORF SEQ. ID NO:	29098		29252		30188	28621	30737	31263	31361	31591		31955	32049			33181		33295		34749	34775		35132			35431	35432	35562	35563			38283
Exen SEQ ID NO:	18626	16696	17106	17247	17757	16139	18265	18538	18626	18820		19158	19246	19780	19863	20284	20437	20683	21197	21799	21826	21874	22160	22160	22304	22449	22449	22567	22567	22610	22798	23248
Probe SEQ ID NO:	4028	4102	4522	4665	5192	5226	5636	5916	9009	6210	6321	0959	9650	7251	7336	7775	7895	8142	8658	9273	9312	9475	9661	9661	9806	9954	9954	10072	10072	10115	10304	10720

Page 92 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

d29807.x1 NCI_CGAP_Ut1 Hamo sapiens cDNA clone IMAGE.2619444 3' similar to gb:M73779 RETINOIC 044h09.s1 Scares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:1526369 3' similar to gb.A03911 Homo sapiens DNA polymerase epsilon catalytic subunit protein (POLE1) gene, exon 1a //42/10.r1 Soares fetal liver spleen 1NFLS Homo sapiens cDNA clone IMAGE:129547 Schizosaccharomyces pombe DNA for cytoplasmic dynein heavy chain, complete cds Pimephales prometas liver glucose-6-phosphate-1-dehydrogenase mRNA, partial cds Homo sapiens lambda/lota protein kinase C-interacting protein mRNA, complete cds Homo sapiens lambda/lota protein kinase C-interacting protein mRNA, complete cds Rattus norvegicus Aryl hydrocarbon receptor nuclear translocator 1 (Arnt1), mRNA ov80a10.s1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:1643610 3 Plasmodium vivax reticulocyte binding protein-2 (rbp-2) gene, complete cds Mus musculus pale ear (ep) gene, wild type allele, 3' region, partial cds Rattus norvegicus chemokine receptor CXCR3 mRNA, complete cds Gallus gallus ovalbumin (Y) gene, complete cds Raftus norvegicus brush border myosin-l (BBMI) mRNA, partial cds RC3-BT0502-251199-011-d01 BT0502 Homo sapiens cDNA Mus musculus interleukin 2 receptor, gamma chain (Il2rg), mRNA GLIA DERIVED NEXIN PRECURSOR (HUMAN); Homo sapiens hypothetical protein FLJ10581 (FLJ10581), mRNA Sigmodon hispidus p53 gene, partial cds Rattus norvegicus arylecetamide deacetylase gene, complete cds Top Hit Descriptor Homo sapiens ninein-Lm isoform (ninein) mRNA, complete cds Homo sapiens Ku70-binding protein (KUB3) mRNA, partial ods Arabidopsis thaliana DNA chromosome 4, contig fragment No. CM3-CT0315-271199-045-b11 CT0315 Homo sapiens cDNA MR1-FN0010-290700-007-d04 FN0010 Hamo sapiens cDNA RC3-BT0502-251189-011-401 BT0502 Homo sapiens cDNA Mus musculus Wm protein (Wrn) gene, complete cds EST67784 Fetal lung II Homo sepiens cDNA 5' end Nouse gene for immunoglobulin diversity region D1 ACID RECEPTOR ALPHA-1 (HUMAN); Sorghum bicolor 22 kDa kafirin cluster EST_HUMAN NT EST_HUMAN EST_HUMAN EST_HUMAN **EST HUMAN** EST_HUMAN EST_HUMAN HUMAN HOMAN Top Hit Database Source EST ST ¥ Ę 8922533 NT 눋 F 7549743 7305180 Top Hit Acession 2.0E-01 AF302773.1 2.0E-01 AW975297.1 2.0E-01 AI023592.1 1.9E-01 AA358813.1 1.9E-01 AW 130149.1 1.9E-01 AF127937.1 2.0E-01 AF078164.2 AF184623.1 BE834943.1 1.9E-01 AF091216.1 1.9E-01 AF004353.1 BE070801.1 BE070801.1 AA916492.1 1.9E-01 AF 264017.1 2.0E-01 D89088.1 2.0E-01 AF206637.2 AF061282.1 AB006784.1 AL161493.2 ş U32581.2 1.9E-01 U66066.1 U25148.1 1.9E-01 D13197. R16467. .9E-01 / 1.9E-01 1.9E-01 1.9E-01 1.9E-01 1.9E-01 1.9E-01 8-01 1.95-01 .9E-01 1.9E-01 .9E-01 1.9E-01 1.9E-01 Vost Simila BLASTE (Top) Hit 5.46 7.81 0.73 1.92 10.04 1.05 4.19 3.85 1.86 5.4 4. 6.6 8.82 4.02 6.58 0.76 69.0 6.22 4 Expression Signal 29123 29202 31173 31360 30788 30888 25509 25792 25793 25800 26256 27331 28128 28939 36264 25800 28039 28528 26534 28611 ORF SEQ ΘNO 18625 13315 13315 13746 14761 15580 15649 16050 16475 16749 18414 18450 24402 24610 13308 13633 14008 14075 15565 SEQ ID 24584 16131 16661 16901 24887 24807 Š 6005 Probe SEQ ID 12162 12449 1023 1023 1143 2422 2949 2965 3033 2828 12374 12425 374 8 3442 3526 4064 4157 4315 4568 5158 5789 69 1413 1482 684 3877 Ö

Page 93 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Single Extri Frobes Expressed III Fetal Liver	. Top Hil Descriptor	AU133116 NT2RP4 Homo sapiens cDNA clone NT2RP4001328 5'	wi34h02.x1 NCI_CGAP_Co16 Homo sapiens cDNA clone IMAGE:23940993'	x14c08.x1 NCI_CGAP_Kid8 Homo sapiens cDNA clone IMAGE:2818030 3' similær to gb:X03559 ATP SYNTHASE BETA CHAIN, MITOCHONDRIAL PRECURSOR (HUMAN);	yg09a12.s1 Soares Infant brain 1NIB Homo sapiens cDNA clone IMAGE:31663 3' similar to contains MER13 repositive element:	Homo saplens tubby like protein 1 (TULP1) gene, exons 9-11	Homo sepiens tubby like protein 1 (TULP1) gene, exons 9-11	Arabidopsis thaliana serina/threonine protein phosphatase type one (TOPP8) gene, complete cds	Zea mays starch branching enzyme I (sbe1) gene, complete cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 57	Homo sapiens mRNA for KIAA1198 protein, partial cds	Marsupial cat beta-globin gene mRNA, partial cds	Marsupial cat beta-globin gene mRNA, partial cds	o98g10.s1 NCI_CGAP_PNS1 Home sapiens cDNA clone IMAGE:1537506 3, similar to contains Alu .	repourte statistic ROS, ETORA, 080700, 022, 402 ETORA, Home, conjens c ONA	RC5-E10082-080700-022-402 FT0082 Homo saniens cDNA	Arabidrosis theliana DNA chromosome 4 contra fragment No. 15	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 15	Homo sepiens calcium channel alpha1E subunit (CACNA1E) gene, exons 7-49, and partial cds, alternatively soliced	Rattus norvegicus sodium channel I mRNA, complete cds	Homo sapiens partial 5-HT4 receptor gene, exons 2 to 5	Drosophila melanogaster clathrin light chain mRNA, complete cds	Arabidopsis thaliana receptor-like kinase LECRK1 (LECRK1) gene, complete cds	Mus musculus p116Rip mRNA, complete cds	Mus musculus Cetg gene for chaperonin containing TCP-1 gamma subunit, partial cds	Homo saplens calcium channel, voltage-dependent, beta 2 subunit (CACNB2) mRNA, and translated	products	Oryzias latipes gene for membrane guanylyl cyclase OIGC1, complete cds	wd71f02.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE:2337051 3'	Dictyostelium discoideum plasmid Ddp5, complete genome	Yersinia pestis plasmid pCD1
LAUI FIUDES	Top Hit Database Source	EST_HUMAN	EST_HUMAN	EST_HUMAN		IN	т			LN		La			EST HIMAN	Т	✝		1			Z	IN		TN				EST_HUMAN	\neg	LN
Alfilio	Top Hit Acession No.	1.9E-01 AU133116.1	1.9E-01 AI762391.1	1.9E-01 AW148452.1	!	=	1.9E-01 AF034920.1	1.9E-01 U80922.1	1.9E-01 AF072724.1		1.1		1.9E-01 M14568.1	1 05 01 0 0012108 1			Τ	1.9E-01 AL161503.2			1.9E-01 AJ243213.1	1.9E-01 AF055900.1	1.9E-01 AF001168.1		1.8E-01 AB022090.1		2532	_			1.8E-01 AL117189.1
	Most Similar (Top) Hit BLAST E Value	1.9E-01	1.9E-01	1.9E-01	1 9F-01 R43212 1	1.85-01/	1.9E-01	1.9E-01	1.9E-01	1.9E-01	1.8E-01	1.9E-01	1.9E-01	1 05 01	10 Hg	1 9E-01	1 BF-01/	1.96-01	1 8F-01 /	1.9E-01	1.9E-01	1.9E-01	1.9E-01	1.8E-01 U73200.1	1.8E-01		1.8E-01	1.8E-01 /	1.8E-01	1.8E-01 /	1.8E-01
	Expression Signal	2.52	1.07	1.23	137	0.91	0.81	1.3	2.89	1.71	12.12	1.36	1.36	27.0	27.0	0.71	200	2.02	2.08	1.68	2.69	1.33	3.69	2.58	1.67		1.76	0.77	0.78	1.26	6.97
	ORF SEQ. ID NO:		31855	31915	30460		32482		15628		34074		34338	14636		l	l	36072	36178		37088			25172	25423						26447
	Exon SEQ ID NO:	18685	19070	19123	18069	19644	19644	20025	20083	20469	21159	21414	21414	22087	22835	22835	23081	23061	23167	23828	24018	24431	24874	12713	15412		13039	13395	13628	13732	13926
	Probe SEQ ID NO:	8048	6469	6253	7050	7072	7072	7503	7543	7927	8820	8875	8875	0820	10140	10140	10523	10523	10835	11377	11571	12207	12582	34	281		38	778	1018	130	1332

Page 94 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Descriptor	Mus musculus guanylate nucleotide binding protein 1 (Gbp1), mRNA	Mus musculus guanylate nucleotide binding protein 1 (Gbp1), mRNA	Homo sapiens latent transforming growth factor beta binding protein 4 (LTBP4) mRNA	qg22dt0.x5 NCI_CGAP_Kid3 Homo sapiens cDNA clone IMAGE:1761811 3' similar to TR:075936 075936 GAMMA BUTYROBETAINE HYDROXYLASE;	Mus musculus Scya6, Scya16-ps, Scya5 genes for small inducible cytokine A6 precursor, small inducible cytokine A5 precursor small cytokine a A5 precursor small cytokine A5 precursor small cyt	QV3-DT0018-081299-036-q04 DT0018 Homo sapiens cDNA	Jonopsidium acaule LEAFY protein (LEAFY2) gene, partial cds	xy41a03.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2659756 3'	QV0-BN0041-070300-147-c04 BN0041 Homo sapiens cDNA	601809723R1 NIH_MGC_18 Homo sapiens cDNA clone IMAGE:4040621 3'	1/45e01.s1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE: 151704 3' similar to contains Alu	repetitive element,	yi45e01.s1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:151704 3' similar to contains Alu repetitive element;	Homo sapiens Xq pseudoautosomal region; segment 1/2	Bowne NB25 mRNA for MHC class II (BcLA-DQB), complete cds	Arabidopsis thaliana DNA chromosome 4, contig fragment No. 56	Mus musculus Scyad, Scyad Scyat 6-ps, Scyad genes for small inducible cytokine A6 precursor, small inducible cytokine A5 precursor, complete cds inducible cytokine A5 precursor, complete cds	S.tuberosum mRNA for alcohol dehydrogenase	MR3-ST0203-151299-112-906 ST0203 Homo sapiens cDNA	an 28g07.y5 Gessler Wilms turnor Homo sapiens cDNA clone IMAGE: 1700028 5'	Mesocricetus auratus Na-taurocholate cotransporting polypeptide mRNA, partial cds	t57e04.x1 NCI_CGAP_Lym12 Homo sapiens cDNA clone IMAGE:2134590 3'	Arabidopsis thaliana cytochrome b-561 (CYTB561) gene, partial cds	Broad bean wilt virus 2 genes encoding 119kDa protein, 104kDa protein, large coat protein, small coat protein	Broad bean wilt virus 2 genes encoding 119kDa protein, 104kDa protein, large coat protein, small coat protein	EST_HUMAN MR4-ST0121-041199-019-b01 ST0121 Homo saplens cDNA
Top Hit Database Source	ĮΣ	Z	Z	EST_HUMAN	<u> </u>	EST HUMAN	IN	EST_HUMAN	EST_HUMAN	EST_HUMAN		EST_HUMAN	EST HUMAN	N	Z	N	IN	IN	EST_HUMAN	EST_HUMAN	IN	EST_HUMAN	TN	IN	LΝ	EST HUMAN
Top Hit Acession No.	6753947 NT	B753947 NT	4505038 NT	AI733708.1	AB051807 1	AW935728.1	AF184589.1	AW182300.1	AW995178.1	BF183582.1		H03369.1	H03369.1	AJ271735.1	D37954.1	AL161556.2	AB051897.1	X92179.1	AW814270.1	A 792382,1	AF181258.1	AI439881.1	AF132115.1	AJ132844.1	AJ132844.1	AW809402.1
Most Similar (Top) Hit BLAST E Value	1.8E-01	1.8E-01	1.8E-01	1.8E-01	10-38	_		1.8E-01	1.8E-01	1.8E-01		1.8E-01	1.8E-01	1.8E-01	1.8E-01	1.8E-01	1.8E-01	1.8E-01			1.8E-01	1.8E-01	1.8E-01	1.8E-01	1.8E-01	1.8E-01
Expression Signal	1.31	1.31	2.79	2.22	1.52	2.29	2.38	1.18	1.31	0.71		0.78	0.79	0.78	4.07	6.59	2.51	1.03	2.18	1.59	1.5	1.07	69'0	0.78	0.78	
ORF SEQ ID NO:		26677			80026			28020		28501		28752	28753			29691	29914	29950		L	30245	30270	30276	30314	30315	
Exan SEQ ID NO:	14143	14143	14472	14492	14542	ł	15540		15772	16021		16284	16284		17012	17236	17461	17503			17820	17843	17850	17899	17899	ΙI
Probe SEQ ID NO:	1551	1551	1887	1907	1058	2716	2923	2928	3158	3413		3683	3683	4333	4426	4654	4886	4928	5198	5216	5257	5281	5288	5338	5338	5398

Page 95 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Top Hit Acession Detabese Top Hit Descriptor Source		4.2 NI	8629.1 EST_HUMAN yx38h08.r1 Soares melanocyte 2NbHM Homo sapiens cDNA clone IMAGE:284083 5"	6678428 NT Mus musculus Trif receptor-essociated factor 6 (Traf6), mRNA	6678428 NT Mus musculus Trif receptor-associated factor 8 (Traf8), mRNA	SWISSPROT	4853.1 EST_HUMAN py62h02.r1 Soares_multiple_sclerosis_2NbHMSP Homo sapiens cDNA chone IMAGE:278163 5'	١	1.8E-01 AB018561.1 NT Citrullus lanatus mRNA for wsus, complete cds	21 BE961353.1 EST_HUMAN 601648361R2 NIH_MGC_62 Homo sepiens cDNA clone IMAGE:3932247 3'	21 AW 966118.1 EST_HUMAN EST378191 MAGE resequences, MAGI Homo saplens cDNA	3258.1 NT Human cellular DNA/Human papillomavirus provinsi DNA	626232	51.1 EST_HUMAN	SWISSPROT	5272 SWISSPROT AMP NUCLEOSIDASE	INT	INT	8123 SWISSPROT COLLAGEN ALPHA 2(I) CHAIN PRECURSOR	7548.1 NT Methanococcus jannaschii section 90 of 150 of the complete genome	!	Z.1 NT	Z	7336.1 NT A. thaitana mRNA for ribonucleotide reductase R2	TN	01 AB018561.1 NT Citrulius lanatus mRNA for wsus, complete cds	IN	01 AF019107.1 NT Dictyostelium discoideum unknown (DG1041) gene, complete cds	IN	7033.1 NT B.teurus mRNA for potassium channel	8394421 NT Rattus norvegicus Thromboxane receptor (Tbxa2r), mRNA	INT	10086561 NT Bovine ephemeral fever virus, complete genome
				6678428 NT	6678428 NT								626232																		8394421 NT		10086561 NT
		1 4 1 1 0	21 N28629.1	01	11	31 Q9QY14	1 N94853.1	1 AB018	1 AB018	1 BE96	1 AW96	1 M732	1.8E-01	1 AA490	01 P15272	21 P15272	D1 M26019.1	01 M26019.1	01 P08123	01 U67548.1		1 AF 200	01 X63440.1	11 X77336.1	01 U38906.1	1 AB01	1 AB01	1 AF018	1.8E-01 M59257.1	1.8E-01 X57033.1	10	-01 040487.1	F
Most Similar (Top) Hit BLAST E Value		1.85-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.85-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0		1.8E-0	1.85-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.8E-0	1.86-0	1.86-0	1.8E	1.8E
Expression Signal			1.01	1.1	1.1	2.03	2.24	1.22	1.22	1.71	0.47	1.13	1.39	0.55	1.13	1.13	0.95	96.0		0.69		0.64	122	2.37	7.47	3.07	3.07	4.49	1.84	4.3	2.74		2.04
ORF SEQ ID NO:		31331	31446	31653	31654	32035		32487	32488	32272	34009	34741	34843		34950	34951	34990	34991	35166	35170			35/53	36066	36106	32487	32488	36160	36434	36045	37111	37132	
Exon SEQ ID NO:	500	18596	18699	18885	18885	19231	19271	19649	19649	19457	21086	21792	21896	21921	21994	21994	22032	22032	22183	22197		22534	22766	23054	23094	19649	19649	23148	23417	23035	24042	24068	24146
Probe SEQ ID NO:	3	9/60	6082	6277	6277	6635	6675	7077	7077	7117	8547	9268	9536	9412	9494	9494	9532	9532	9694	8698		10039	10271	10518	10558	10615	10615	10616	10897	11337	11599	11626	11748

Page 96 of 526
Table 4
Single Exon Probes Expressed in Fetal Liver

Homo sapiens derivative 11 breakpoint fragment; partial intron 10 of the ALL-1/ML/HRX gene fused to intron qh57e09.x1 Soares_fetal_liver_splean_1NFLS_S1 Homo sapiens cDNA clone IMAGE:1848808 3' similar to Anabaena sp. ORF4 (partial), ORF3, ORF2, ORF1, adpA gene, adpB gene, adpC gene, adpD gene, adpE Vibrio chalerae hypoxanthine phosphoribosyltransferase (hpt) gene, partial cds, hemagglutinin/protease Vibrio cholerae hypoxanthine phosphoribosyltransferase (hpt) gene, partial cds, hemagglutinin/protease P. dumerilii histone gene cluster for core histones H2A, H2B, H3 and H4 NEUROFILAMENT TRIPLET L PROTEIN (NEUROFILAMENT LIGHT POLYPEPTIDE) (NF-L) Lymantria dispar nucleopolyhedrovirus, complete genome
Arabidopsis thaliana DNA chromosome 4, contig fragment No. 69
Homo sapiens BNIP3H (BNIP3H) gene, complete cds; nuclear gene for mitochondrial product 12346F Human fetal heart, Lambda ZAP Express Homo sapiens cDNA clone J2346 5 yh48h10.r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:133027 5' Homo sapiens fragile 16D oxido reductase (FOR) gene, exons 8, 9, and partial cds regulatory protein (hapR) gene, complete cds, and YRAL VIBCO gene, partial cds Brn67 Homo sapiens cDNA clone IMAGE:4155318 5' regulatory protein (hapR) gene, complete cds, and YRAL VIBCO gene, partial cds Zea mays calcium-dependent protein kinase (MZECDPK2) mRNA, complete cds Rattus norvegicus procollegen C-proteinase enhancer protein (Pcolce), mRNA exus canadensis gerany/gerany diphosphate synthase mRNA, complete cds 602186630F1 NIH_MGC_49 Homo sepiens cDNA clone IMAGE:4298646 5 601274604F1 NIH_MGC_20 Homo sepiens cDNA clone IMAGE:3615768 5 DNA TERMINAL PROTEIN (BELLETT PROTEIN) (PTP PROTEIN) Fop Hit Descriptor EST41651 Endometrial tumor Homo sapiens cDNA 5' end Lymentria disper nucleopolyhedrovirus, complete genome Schistocerca gregaria alpha repetitive DNA contains OFR.b1 OFR repetitive element E.dispar mRNA for hexokinase (hxk1) Naja naja atra ctx-1 gene, exons 1-3 Naja naja atra cbx-1 gene, exons 1-3 Yersinia pestis plasmid pCD 5 of the AF-4/FEL gene gene and adpF gene 502019928F1 NCI SWISSPROT EST_HUMAN EST_HUMAN NT EST HUMAN EST_HUMAN EST_HUMAN EST_HUMAN HUMAN Top Hit Database Source SWISSPR01 RST. 눌 Ę 뉟뉟 눋 þ ź 닐닏 ż Ę 'n ١ F Top Hit Acession 1.7E-01 Al247635.1 1.7E-01 U28376.1 1.7E-01 BF689719.1 AL161573.2 AF255051.1 BE385164.1 1.7E-01 P35616 1.7E-01 AF081810.1 1.7E-01 AJ238736.1 1.7E-01 AJ238736.1 1.7E-01 AA336909.1 1.7E-01 AF081514.1 1.7E-01 N55763.1 AF217490.1 AL117189.1 1.7E-01 AF081810,1 AF000716.1 AF000716.1 AJ269505.1 1.7E-01 AJ235377.1 ģ 1.7E-01 X52936.1 1.7E-01 AF217490 1.7E-01 X53330.1 R24494.1 1.8E-01 Y11114.1 1.8E-01 Q96682 1.8E-01 1.7E-01 1.7E-01 1.8E-01 1.8E-01 1.7E-01 1.7E-01 1.7E-01 1.8E-01 Aost Similar (Top) Hit BLAST E Value 8 8 5.53 8.63 0.67 1.98 1.98 1.53 1.9 1.91 1,3 8 1.07 1.02 1.58 2.99 0.67 0.95 5.61 Expression Signal 27973 28121 30227 30507 26210 27974 28220 28574 29083 26447 25964 26211 2804 29904 29981 ORF SEQ 31032 26998 28500 ΘNΟ 15643 15643 25045 15503 15503 18610 17240 17539 17806 24186 24569 24590 13232 13454 13608 13701 15569 15753 16099 17775 SEQ ID 14441 16020 17452 24491 Š 5242 4965 1096 2025 2953 3494 4012 Probe SEQ ID 12218 12416 12459 12502 3139 4658 4877 11814 12291 603 838 966 2885 2885 3027 3027 3412 ö

Page 97 of 526 Table 4 Single Exon Probes Expressed in Fetal Liver

Most Similar (Top) Hit Acession BLAST E No. Value 1.7E-01 AF072725.1
1.7E-01 J04479.1 NT
1.7E-01 AA470686.1 EST_HUMAN
1.7E-01 AA470686.1 EST_HUMAN
1.7E-01 U43599.1
H72118.1
AI370976.1
AI370976.1
BE300286.1
AF026552.3
292910.1
AP000422.1
1.7E-01 BE734179.1 EST HUMAN 1.7E-01 P16724 SWISSPROT
Q01955
AF000573.1
AF15066
8428
3.1
1.7E-01 BE253142.1 EST_HUMAN
1.7E-01 AP001508.1 NT
1.7E-01 AW977455.1 EST_HUMAN
1.7E-01[AW977455.1 [EST_HUMAN
1.7E-01 U16288.1 NT
1.7E-01 Z34508.1 NT